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OM protein - protein search, using sw model

Run on: May 4, 2005, 18:14:26 ; Search time 134 Seconds
(without alignments)
116.836 Million cell updates/sec

Title: US-10-723-368-5

Perfect score: 254
Sequence: 1 TQQLRVGCVLGTCTGVQNL.....MGPAGRQDSAPVDPSSPHSY 47

Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 1426032 seqs, 333106140 residues

Total number of hits satisfying chosen parameters: 1426032

Minimum DB seq length: 0
Maximum DB seq length: 200000000

Post-processing: Minimum Match 100%
Maximum Match 100%

Listing first 45 summaries

Database :

Published Applications AA:*

- 1: /cgn2_6/prodata/1/pubppaa/US07_PUBCOMB.pep:*
- 2: /cgn2_6/prodata/1/pubppaa/PCR_NEW_PUB.pep:*
- 3: /cgn2_6/prodata/1/pubppaa/US06_NEW_PUB.pep:*
- 4: /cgn2_6/prodata/1/pubppaa/US06_PUBCOMB.pep:*
- 5: /cgn2_6/prodata/1/pubppaa/US07_NEW_PUB.pep:*
- 6: /cgn2_6/prodata/1/pubppaa/PCRUS_PUBCOMB.pep:*
- 7: /cgn2_6/prodata/1/pubppaa/US08_NEW_PUB.pep:*
- 8: /cgn2_6/prodata/1/pubppaa/US08_PUBCOMB.pep:*
- 9: /cgn2_6/prodata/1/pubppaa/US09_PUBCOMB.pep:*
- 10: /cgn2_6/prodata/1/pubppaa/US09B_PUBCOMB.pep:*
- 11: /cgn2_6/prodata/1/pubppaa/US09C_PUBCOMB.pep:*
- 12: /cgn2_6/prodata/1/pubppaa/US09_NEW_PUB.pep:*
- 13: /cgn2_6/prodata/1/pubppaa/US10A_PUBCOMB.pep:*
- 14: /cgn2_6/prodata/1/pubppaa/US10B_PUBCOMB.pep:*
- 15: /cgn2_6/prodata/1/pubppaa/US10C_PUBCOMB.pep:*
- 16: /cgn2_6/prodata/1/pubppaa/US10D_PUBCOMB.pep:*
- 17: /cgn2_6/prodata/1/pubppaa/US10E_NEW_PUB.pep:*
- 18: /cgn2_6/prodata/1/pubppaa/US11_NEW_PUB.pep:*
- 19: /cgn2_6/prodata/1/pubppaa/US60_NEW_PUB.pep:*
- 20: /cgn2_6/prodata/1/pubppaa/US60_PUBCOMB.pep:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	254	100.0	47	17	US-10-850-055-14
2	72.5	28.5	52	9	US-09-813-345-14
3	72.5	28.5	52	14	US-10-197-954-2
4	72.5	28.5	52	16	US-10-474-635A-19
5	72.5	28.5	52	17	US-10-718-071-43
6	72.5	28.5	52	17	US-10-760-085-2
7	72.5	28.5	185	15	US-10-364-889-6
8	72.5	28.5	185	15	US-10-372-683-12
9	72.5	28.5	185	16	US-10-675-406A-7
10	72.5	28.5	185	16	US-10-755-889-148
11	69.5	27.4	50	9	US-09-813-345-15
12	68.5	27.0	52	17	US-10-718-071-16
13	64	25.2	103	15	US-10-264-237-1622

14	63.5	25.0	216	16	US-10-437-963-198108	Sequence 198108,
15	62.5	24.6	52	15	US-10-360-101-74	Sequence 74, App1
16	59	23.2	142	15	US-10-307-817-14	Sequence 14, App1
17	58.5	23.0	92	16	US-10-437-963-198935	Sequence 198935,
18	58.5	23.0	521	16	US-10-437-963-124602	Sequence 26, App1
19	58.5	23.0	576	15	US-10-184-648-26	Sequence 8, App1
20	58	22.8	560	14	US-10-238-129-8	Sequence 8, App1
21	58	22.8	560	14	US-10-238-667-8	Sequence 6078, Ap
22	57	22.8	1544	15	US-10-369-493-6078	Sequence 6078, Ap
23	57	22.4	117	16	US-10-767-701-51114	Sequence 51114, A
24	56.5	22.2	575	15	US-10-425-114-64569	Sequence 64569, A
25	56	22.0	89	16	US-10-437-963-136207	Sequence 136207,
26	56	22.0	107	9	US-09-764-870-356	Sequence 356, App
27	56	22.0	107	9	US-09-764-853-470	Sequence 470, App
28	56	22.0	107	14	US-10-125-540-356	Sequence 356, App
29	56	22.0	107	14	US-10-103-313-301	Sequence 301, App
30	56	22.0	107	15	US-10-158-057-335	Sequence 335, App
31	56	22.0	947	15	US-10-461-194-114	Sequence 114, App
32	55	21.7	220	16	US-10-767-701-32051	Sequence 32051, A
33	55	21.7	322	15	US-10-424-599-175411	Sequence 175411,
34	55	21.7	669	15	US-10-120-801-108	Sequence 108, App
35	54.5	21.5	126	16	US-10-437-963-202043	Sequence 202043,
36	54.5	21.5	320	16	US-10-437-963-119917	Sequence 119917,
37	54.5	21.5	751	16	US-10-437-963-135754	Sequence 135754,
38	54	21.3	538	16	US-10-437-963-151317	Sequence 151317,
39	53.5	21.1	135	15	US-10-264-049-4093	Sequence 4093, Ap
40	53.5	21.1	303	14	US-10-097-340-143	Sequence 143, App
41	53.5	21.1	303	14	US-10-171-311-87	Sequence 87, App1
42	53.5	21.1	329	10	US-09-999-121-6	Sequence 48541, A
43	53	20.9	86	9	US-09-864-761-48541	Sequence 41, App1
44	53	20.9	117	10	US-09-789-390-41	Sequence 8699, Ap
45	53	20.9	136	14	US-10-156-761-8659	

ALIGNMENTS

RESULT 1
US-10-850-055-14
Sequence 14, Application US/10850055
Publication No. US2005009742A1
GENERAL INFORMATION:
APPLICANT: Bertilsson, Goran
APPLICANT: Fritsen, Jonas
APPLICANT: Haegerstrand, Anders
APPLICANT: Heidrich, Jessica
APPLICANT: Hellstrom, Kristina
APPLICANT: Hagbladh, Johan
APPLICANT: Jansson, Katarina
APPLICANT: Kortessan, Jarkko
APPLICANT: Lindquist, Per
APPLICANT: Lundh, Hanna
APPLICANT: McGuire, Jacqueline
APPLICANT: Mercer, Alex
APPLICANT: Nyberg, Karl
APPLICANT: Oseolnak, Amna
APPLICANT: Patrone, Cesare
APPLICANT: Ronholm, Harriet
APPLICANT: Wikstrom, Lillian
APPLICANT: Zachrisson, Olof
TITLE OF INVENTION: COMPOUNDS AND METHODS FOR INCREASING NEUROGENESIS
FILE REFERENCE: 21882-517 CIP
CURRENT APPLICATION NUMBER: US/10/850,055
CURRENT FILING DATE: 2004-05-19
PRIOR APPLICATION NUMBER: US 10/718,071
PRIOR FILING DATE: 2003-11-20
PRIOR APPLICATION NUMBER: US 60/427,912
PRIOR FILING DATE: 2002-11-20
NUMBER OF SEQ ID NOS: 66
SOFTWARE: PatentIn version 3.2
SEQ ID NO 14
LENGTH: 47

TYPE: PRT
ORGANISM: Homo sapiens
US-10-850-035-14

Query Match 100.0%; Score 254; DB 17; Length 47;
Best Local Similarity 100.0%; Pred. No. 1.2e-25;
Matches 47; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TQQLRVGCVLTGCOYONSHRLMQLMGPAGQDSAPVDPSSPHSY 47
DB 1 TQQLRVGCVLTGCOYONSHRLMQLMGPAGQDSAPVDPSSPHSY 47

RESULT 2

US-09-813-345-14
Sequence 14, Application US/09813345
Patent No. US20020068814A1
GENERAL INFORMATION:

APPLICANT: Smith, Derek D.
Saha, Shankar
Abel, Peter W.

TITLE OF INVENTION: PEPTIDE ANTAGONISTS OF GCRP-RECEPTOR
SUPERFAMILY AND METHODS OF USE

NUMBER OF SEQUENCES: 23

CORRESPONDENCE ADDRESS:
ADDRESSEE: Mueling, Raasch & Gebhardt, P.A.
STREET: 119 No. US20020068814A1th Fourth Street
CITY: Minneapolis

STATE: MN

COUNTRY: USA

ZIP: 55401

COMPUTER READABLE FORM:

MEDIUM TYPE: Floppy disk

OPERATING SYSTEM: PC-DOS/MS-DOS

SOFTWARE: Patent in Release #1.0, Version #1.30

CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/813,345
FILING DATE: 20-Mar-2001

CLASSIFICATION: <Unknown>

ATTORNEY/AGENT INFORMATION:

NAME: McCormick, Myra H

REGISTRATION NUMBER: 36,602

REFERENCE/DOCKET NUMBER: 180.00020101

TELECOMMUNICATION INFORMATION:
TELEPHONE: 612/305-1228
TELEFAX: 612/305-1228

INFORMATION FOR SEQ ID NO: 14:

SEQUENCE CHARACTERISTICS:

LENGTH: 52 amino acids

TYPE: amino acid

STRANDEDNESS: single

TOPOLOGY: linear

MOLECULE TYPE: peptide

SEQUENCE DESCRIPTION: SEQ ID NO: 14:

US-09-813-345-14

Query Match 28.5%; Score 72.5; DB 9; Length 52;

Best Local Similarity 38.5%; Pred. No. 0.067; 17; Indels 1; Gaps 1;

Matches 15; Conservative 6; Mismatches 17; Indels 1; Gaps 1;

QY 9 GCVLTGCOYONSHRLMQLMGPAGQDSAPVDPSSPHSY 47

DB 15 GCRFGCTVQKLAHQIYQFT-DKDKDNVAPRSKISPGY 52

RESULT 3

US-10-197-954-2

Sequence 2, Application US/10197954

Publication No. US20030119021A1

GENERAL INFORMATION:

APPLICANT: Krater, Hubert

APPLICANT: Siddiqui, Suhail

APPLICANT: Little, Daniel
TITLE OF INVENTION: Capture Compounds, Collections Thereof
TITLE OF INVENTION: And Methods For Analyzing The Proteome And Complex
TITLE OF INVENTION: Compositions

FILE REFERENCE: 24743-2305

CURRENT APPLICATION NUMBER: US/10/197,954

CURRENT FILING DATE: 2002-07-16

PRIOR APPLICATION NUMBER: 60/306,019

PRIOR FILING DATE: 2001-07-16

PRIOR APPLICATION NUMBER: 60/314,123

PRIOR FILING DATE: 2001-08-21

PRIOR APPLICATION NUMBER: 60/363,433

PRIOR FILING DATE: 2002-03-11

NUMBER OF SEQ ID NOS: 149

SOFTWARE: FastSeq for Windows Version 4.0

SEQ ID NO 2

LENGTH: 52

TYPE: PRT

ORGANISM: Homo Sapien

US-10-197-954-2

Query Match 28.5%; Score 72.5; DB 14; Length 52;

Best Local Similarity 38.5%; Pred. No. 0.067; 17; Indels 1; Gaps 1;

Matches 15; Conservative 6; Mismatches 17; Indels 1; Gaps 1;

QY 9 GCVLTGCOYONSHRLMQLMGPAGQDSAPVDPSSPHSY 47

DB 15 GCRFGCTVQKLAHQIYQFT-DKDKDNVAPRSKISPGY 52

RESULT 4

US-10-474-635A-19

Sequence 19, Application US/10474635A

Publication No. US20040176567A1

GENERAL INFORMATION:

APPLICANT: Isis Innovation Ltd

TITLE OF INVENTION: Peptides

FILE REFERENCE: 480821.00004

CURRENT APPLICATION NUMBER: US/10/474,635A

CURRENT FILING DATE: 2003-10-14

PRIOR APPLICATION NUMBER: GB 0109438.2

PRIOR FILING DATE: 2001-04-17

NUMBER OF SEQ ID NOS: 22

SOFTWARE: Patent in version 3.1

SEQ ID NO 19

LENGTH: 52

TYPE: PRT

ORGANISM: Homo sapiens

US-10-474-635A-19

Query Match 28.5%; Score 72.5; DB 16; Length 52;

Best Local Similarity 38.5%; Pred. No. 0.067; 17; Indels 1; Gaps 1;

Matches 15; Conservative 6; Mismatches 17; Indels 1; Gaps 1;

QY 9 GCVLTGCOYONSHRLMQLMGPAGQDSAPVDPSSPHSY 47

DB 15 GCRFGCTVQKLAHQIYQFT-DKDKDNVAPRSKISPGY 52

RESULT 5

US-10-718-071-43

Sequence 43, Application US/10718071

Publication No. US20050009847A1

GENERAL INFORMATION:

APPLICANT: Bertilsson, Goran

APPLICANT: Eriandsson, Rikard

APPLICANT: Filsen, Jonas

APPLICANT: Haegerstrand, Anders

APPLICANT: Heldric, Jessica

APPLICANT: Hellstrom, Kristina

APPLICANT: Haggblad, Johan

APPLICANT: Jansson, Katarina

APPLICANT: Korlesmaa, Jarkko

APPLICANT: Lindquist, Per
APPLICANT: Lundh, Hanna
APPLICANT: McGuire, Jacqueline
APPLICANT: Mercer, Alex
APPLICANT: Nyberg, Karl
APPLICANT: Ossolink, Amlna
APPLICANT: Patrone, Cesare
APPLICANT: Ronholm, Harriet
APPLICANT: Wikstrom, Lillian
APPLICANT: Zachrisson, Olof
TITLE OF INVENTION: COMPOUNDS AND METHODS FOR INCREASING NEUROGENESIS
FILE REFERENCE: 21882-517 UTIL
CURRENT APPLICATION NUMBER: US/10/718,071
CURRENT FILING DATE: 2003-11-20
PRIOR APPLICATION NUMBER: US 60/427,912
PRIOR FILING DATE: 2002-11-20
NUMBER OF SEQ ID NOS: 71
SOFTWARE: PatentIn version 3.2
SEQ ID NO 43
LENGTH: 52
TYPE: PRT
ORGANISM: Homo sapiens
US-10-718-071-43

Query Match 28.5%; Score 72.5; DB 17; Length 52;
Best Local Similarity 38.5%; Pred. No. 0.067;
Matches 15; Conservative 6; Mismatches 17; Indels 1; Gaps 1;

QY 9 GCVLGTCOVONTLSHRLMQLMGAPAGRODSAPVDPSSPHSY 47
DB 15 GCRFGTCVTQKLAHQIYQFT-DKDKDNVAPRSKISPGY 52

RESULT 6
US-10-760-085-2
Sequence 2, Application US/10760085
Publication No. US20050042771A1
GENERAL INFORMATION:
APPLICANT: Hubert K*ster
APPLICANT: Daniel Paul Little
APPLICANT: Suhaid Mahmood Siddiqi
APPLICANT: Matlew Peter Grealish
APPLICANT: Subramaniam Mareppan
APPLICANT: Chester Frederick Haseman III
APPLICANT: Ping Yip
TITLE OF INVENTION: Capture Compounds, Collections Thereof
TITLE OF INVENTION: And Methods For Analyzing The Proteome And Complex
FILE REFERENCE: 24743-2309
CURRENT APPLICATION NUMBER: US/10/760,085
CURRENT FILING DATE: 2004-01-16
PRIOR APPLICATION NUMBER: 60/441,398
PRIOR FILING DATE: 2003-01-16
NUMBER OF SEQ ID NOS: 149
SOFTWARE: FaastSeq for Windows Version 4.0
SEQ ID NO 2
LENGTH: 52
TYPE: PRT
ORGANISM: Homo Sapien
US-10-760-085-2

Query Match 28.5%; Score 72.5; DB 17; Length 52;
Best Local Similarity 38.5%; Pred. No. 0.067;
Matches 15; Conservative 6; Mismatches 17; Indels 1; Gaps 1;

QY 9 GCVLGTCOVONTLSHRLMQLMGAPAGRODSAPVDPSSPHSY 47
DB 15 GCRFGTCVTQKLAHQIYQFT-DKDKDNVAPRSKISPGY 52

RESULT 7
US-10-364-889-6
Sequence 6, Application US/10364889

Publication No. US20030224989A1
GENERAL INFORMATION:
APPLICANT: Pabel, Gregory L.
APPLICANT: Quinn, Kerry
TITLE OF INVENTION: Compositions and Methods for Treatment of Osteoarthritis
FILE REFERENCE: 21402-558
CURRENT APPLICATION NUMBER: US/10/364,889
CURRENT FILING DATE: 2003-02-12
PRIOR APPLICATION NUMBER: 60/356,376
PRIOR FILING DATE: 2002-02-12
NUMBER OF SEQ ID NOS: 8
SOFTWARE: CuraseqList version 0.1
SEQ ID NO 6
LENGTH: 185
TYPE: PRT
ORGANISM: Homo sapiens
US-10-364-889-6

Query Match 28.5%; Score 72.5; DB 15; Length 185;
Best Local Similarity 38.5%; Pred. No. 0.27;
Matches 15; Conservative 6; Mismatches 17; Indels 1; Gaps 1;

QY 9 GCVLGTCOVONTLSHRLMQLMGAPAGRODSAPVDPSSPHSY 47
DB 109 GCRFGTCVTQKLAHQIYQFT-DKDKDNVAPRSKISPGY 146

RESULT 8
US-10-372-683-12
Sequence 12, Application US/10372683
Publication No. US20040009171A1
GENERAL INFORMATION:
APPLICANT: GERRITSEN, MARY E.
APPLICANT: PERLS JR., FRANKLIN V.
APPLICANT: WU, THOMAS D.
TITLE OF INVENTION: METHODS FOR THE TREATMENT OF CARCINOMA
FILE REFERENCE: P1928R1P1
CURRENT APPLICATION NUMBER: US/10/372,683
CURRENT FILING DATE: 2003-02-21
PRIOR APPLICATION NUMBER: US 10/271,690
PRIOR FILING DATE: 2002-10-16
PRIOR APPLICATION NUMBER: US 60/344,534
PRIOR FILING DATE: 2001-10-18
NUMBER OF SEQ ID NOS: 49
SEQ ID NO 12
LENGTH: 185
TYPE: PRT
ORGANISM: Homo sapien
US-10-372-683-12

Query Match 28.5%; Score 72.5; DB 15; Length 185;
Best Local Similarity 38.5%; Pred. No. 0.27;
Matches 15; Conservative 6; Mismatches 17; Indels 1; Gaps 1;

QY 9 GCVLGTCOVONTLSHRLMQLMGAPAGRODSAPVDPSSPHSY 47
DB 109 GCRFGTCVTQKLAHQIYQFT-DKDKDNVAPRSKISPGY 146

RESULT 9
US-10-675-406A-7
Sequence 7, Application US/10675406A
Publication No. US20040121375A1
GENERAL INFORMATION:
APPLICANT: Bayer Pharmaceuticals Corporation
APPLICANT: Eweleigh, Deepra
APPLICANT: Taylor, Ian
TITLE OF INVENTION: METHODS FOR PREDICTION AND PROGNOSIS OF CANCER, AND MONITORING
FILE REFERENCE: 5138
CURRENT APPLICATION NUMBER: US/10/675,406A
CURRENT FILING DATE: 2003-09-30
PRIOR APPLICATION NUMBER: US 60/415,194

PRIOR FILING DATE: 2002-09-30
NUMBER OF SEQ ID NOS: 7
SOFTWARE: Patentin version 3.2
SEQ ID NO 7
LENGTH: 185
TYPE: PRT
ORGANISM: Homo sapiens
US-10-675-406A-7

Query Match 28.5%; Score 72.5; DB 16; Length 185;
Best Local Similarity 38.5%; Pred. No. 0.27;
Matches 15; Conservative 6; Mismatches 17; Indels 1; Gaps 1;

QY 9 GCVLGTCOVONLSHRLMQLMGAPAGRODSAPVDPSSPHSY 47
DB 109 GCRFGCTVQKLAHQIYQFT-DKDKXNVAFRSKISFGY 146

RESULT 10
US-10-755-889-148
Sequence 148, Application US/10755889
Publication No. US20040171823A1
GENERAL INFORMATION:
APPLICANT: Bristol-Myers Squibb Company
TITLE OF INVENTION: POLYNUCLEOTIDES AND POLYPEPTIDES ASSOCIATED WITH THE NF-KB
FILE REFERENCE: D0284 NP
CURRENT APPLICATION NUMBER: US/10/755,889
PRIOR FILING DATE: 2004-01-13
PRIOR APPLICATION NUMBER: U.S. 60/440,068
PRIOR FILING DATE: 2003-01-14
PRIOR APPLICATION NUMBER: U.S. 60/469,757
NUMBER OF SEQ ID NOS: 823
SOFTWARE: Patentin version 3.2
SEQ ID NO 148
LENGTH: 185
TYPE: PRT
ORGANISM: Homo sapiens
US-10-755-889-148

* Query Match 28.5%; Score 72.5; DB 16; Length 185;
Best Local Similarity 38.5%; Pred. No. 0.27;
Matches 15; Conservative 6; Mismatches 17; Indels 1; Gaps 1;

QY 9 GCVLGTCOVONLSHRLMQLMGAPAGRODSAPVDPSSPHSY 47
DB 109 GCRFGCTVQKLAHQIYQFT-DKDKXNVAFRSKISFGY 146

RESULT 11
US-09-813-345-15
Sequence 15, Application US/09813345
Patent No. US20020068814A1
GENERAL INFORMATION:
APPLICANT: Smith, Derek D.
Sana, Shankar
Abel, Peter W.
TITLE OF INVENTION: PEPTIDE ANTAGONISTS OF CGRP-RECEPTOR
SUPERFAMILY AND METHODS OF USE
NUMBER OF SEQUENCES: 23
CORRESPONDENCE ADDRESS:
ADDRESSEE: Muehling, Raasch & Gebhardt, P.A.
STREET: 119 No. US20020068814A1ch Fourth Street
CITY: Minneapolis
STATE: MN
COUNTRY: USA
ZIP: 55401
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.30

CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/813,345
FILING DATE: 20-Mar-2001
CLASSIFICATION: <Unknown>
ATTORNEY/AGENT INFORMATION:
NAME: McCormack, Myra H
REGISTRATION NUMBER: 36,602
REFERENCE/DOCKET NUMBER: 180.00020101
TELECOMMUNICATION INFORMATION:
TELEPHONE: 612/305-1220
TELEFAX: 612/305-1228
INFORMATION FOR SEQ ID NO: 15:
SEQUENCE CHARACTERISTICS:
LENGTH: 50 amino acids
TYPE: amino acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: peptide
SEQUENCE DESCRIPTION: SEQ ID NO: 15:
US-09-813-345-15

Query Match 27.4%; Score 69.5; DB 9; Length 50;
Best Local Similarity 35.9%; Pred. No. 0.16;
Matches 14; Conservative 7; Mismatches 17; Indels 1; Gaps 1;

QY 9 GCVLGTCOVONLSHRLMQLMGAPAGRODSAPVDPSSPHSY 47
DB 13 GCRFGCTVQKLAHQIYQFT-DKDKXNVAFRSKISFGY 50

RESULT 12
US-10-718-071-16
Sequence 16, Application US/10718071
Publication No. US2005009847A1
GENERAL INFORMATION:
APPLICANT: Bertilsson, Goran
APPLICANT: Erlandsson, Rikard
APPLICANT: Friesen, Jonas
APPLICANT: Haegerstrand, Anders
APPLICANT: Heidrich, Uesastica
APPLICANT: Hellstrom, Kristina
APPLICANT: Hagblad, Johan
APPLICANT: Jansson, Katarina
APPLICANT: Kortessmaa, Jarkko
APPLICANT: Lindquist, Per
APPLICANT: Lundh, Hanna
APPLICANT: McGuire, Jacqueline
APPLICANT: Mercer, Alex
APPLICANT: Nyberg, Karl
APPLICANT: Ossolinak, Amina
APPLICANT: Patrone, Cesare
APPLICANT: Ronnholm, Harriet
APPLICANT: Wikstrom, Lilian
APPLICANT: Zachrisson, Olof
TITLE OF INVENTION: COMPOUNDS AND METHODS FOR INCREASING NEUROGENESIS
FILE REFERENCE: 21882-517 UTIL
CURRENT APPLICATION NUMBER: US/10/718,071
PRIOR FILING DATE: 2003-11-20
PRIOR APPLICATION NUMBER: US 60/427,912
PRIOR FILING DATE: 2002-11-20
NUMBER OF SEQ ID NOS: 71
SOFTWARE: Patentin version 3.2
SEQ ID NO 16
LENGTH: 52
TYPE: PRT
ORGANISM: Homo sapiens
US-10-718-071-16

Query Match 27.0%; Score 68.5; DB 17; Length 52;
Best Local Similarity 38.5%; Pred. No. 0.22;
Matches 15; Conservative 5; Mismatches 18; Indels 1; Gaps 1;

QY 9 GCVLGTCOVONLSHRLMQLMGAPAGRODSAPVDPSSPHSY 47

Db 15 GCRFGTCTVQKLAHQIYQFT-DKDKONVAPRSKISFGY 52

RESULT 13

US-10-264-237-1622
; Sequence 1622, Application US/10264237
; Publication No. US20040009491A1
; GENERAL INFORMATION:
; APPLICANT: Birse et al.
; TITLE OF INVENTION: Nucleic Acids, Proteins, and Antibodies
; FILE REFERENCE: PA131P1
; CURRENT APPLICATION NUMBER: US/10/264,237
; CURRENT FILING DATE: 2002-10-04
; PRIOR APPLICATION NUMBER: PCT/US01/16450
; PRIOR FILING DATE: 2001-05-18
; PRIOR APPLICATION NUMBER: US 60/205,515
; PRIOR FILING DATE: 2000-05-19
; NUMBER OF SEQ ID NOS: 2876
; SOFTWARE: PatentIn Ver. 3.1
; SEQ ID NO 1622
; LENGTH: 103
; TYPE: PRT
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: MISC_FEATURE
; LOCATION: (62)
; OTHER INFORMATION: Xaa equals any of the twenty naturally occurring L-amino acids
; FEATURE:
; NAME/KEY: MISC_FEATURE
; LOCATION: (67)
; OTHER INFORMATION: Xaa equals any of the twenty naturally occurring L-amino acids
; NAME/KEY: MISC_FEATURE
; LOCATION: (71)
; OTHER INFORMATION: Xaa equals any of the twenty naturally occurring L-amino acids
; FEATURE:
; NAME/KEY: MISC_FEATURE
; LOCATION: (102)
; OTHER INFORMATION: Xaa equals any of the twenty naturally occurring L-amino acids
US-10-264-237-1622
Query Match 25.2%; Score 64; DB 15; Length 103;
Best Local Similarity 40.0%; Pred. No. 1.8;
Matches 18; Conservative 5; Mismatches 14; Indels 8; Gaps 3;
QY 2 QAOQLRVGCVLGTCCQVONLSHRLWQMGPAGRDSDAPVDPSSPHS 46
Db 8 QAOQLDSDGC-QQTSPLDPDNNHNAW-ILGPPS-----LDPRSPTS 44
RESULT 14
US-10-437-963-198108
; Sequence 198108, Application US/10437963
; Publication No. US20040123343A1
; GENERAL INFORMATION:
; APPLICANT: La Rosa, Thomas J.
; APPLICANT: Kovacic, David K.
; APPLICANT: Zhou, Yihua
; APPLICANT: Cao, Yongwei
; APPLICANT: Wu, Wei
; APPLICANT: Boukharov, Andrey A.
; APPLICANT: Barabak, Brad
; APPLICANT: Li, Ping
; TITLE OF INVENTION: Rice Nucleic Acid Molecules and Other Molecules Associated With
; TITLE OF INVENTION: Plants and Uses Thereof for Plant Improvement
; FILE REFERENCE: 38-21(53221)B
; CURRENT APPLICATION NUMBER: US/10/437,963
; CURRENT FILING DATE: 2003-05-14
; NUMBER OF SEQ ID NOS: 204966
; SEQ ID NO 198108
; LENGTH: 216
; TYPE: PRT

; ORGANISM: Oryza sativa
; FEATURE:
; OTHER INFORMATION: Clone ID: PAT_MRT4530_93800C.1.pcp
US-10-437-963-198108

Query Match 25.0%; Score 63.5; DB 16; Length 216;
Best Local Similarity 31.0%; Pred. No. 4.7;
Matches 13; Conservative 13; Mismatches 11; Indels 5; Gaps 1;

QY 6 LRVGCVLGTCCQVONLSHRLWQMGPAGRDSDAPVDPSSPHS 42
Db 19 LGRGIIIGTCVTSVSNRBLGFAVNRVHGSGSKQTAEIDPS 60

RESULT 15

US-10-360-101-74
; Sequence 74, Application US/10360101
; Publication No. US20040009550A1
; GENERAL INFORMATION:
; APPLICANT: Moill, Gert N.
; APPLICANT: Leenhouts, Cornelis J.
; TITLE OF INVENTION: Export and modification of (poly)peptide in the lantibiotic way
; FILE REFERENCE: 2183-5673
; CURRENT APPLICATION NUMBER: US/10/360,101
; CURRENT FILING DATE: 2003-02-07
; PRIOR APPLICATION NUMBER: EP 02077060.8
; PRIOR FILING DATE: 2002-05-24
; NUMBER OF SEQ ID NOS: 309
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 74
; LENGTH: 52
; TYPE: PRT
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: A13,S16-sequence of Adrenomedullin Hypotensive peptide
US-10-360-101-74

Query Match 24.6%; Score 62.5; DB 15; Length 52;
Best Local Similarity 35.9%; Pred. No. 1.4;
Matches 14; Conservative 6; Mismatches 18; Indels 1; Gaps 1;

QY 9 GCVLGTCCQVONLSHRLWQMGPAGRDSDAPVDPSSPHS 47
Db 15 GSRFGTCTVQKLAHQIYQFT-DKDKONVAPRSKISFGY 52

Search completed: May 4, 2005, 18:27:15
Job time: 135 secs

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GenCore version 5.1.6
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OM protein - protein search, using sw model

Run on: May 4, 2005, 17:56:39 ; Search time 162 Seconds
(without alignments)
112.208 Million cell updates/sec

Title: US-10-723-368-5
Perfect score: 254
Sequence: 1 TQAQLRVGCVLTGTCVQNTL.....MGPAGRODSAPVDPSSPHSY 47

Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 2105692 seqs, 386760381 residues

Total number of hits satisfying chosen parameters: 2105692

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Database : A_Geneseq_16Dec04:*

- 1: geneseqp1980s:*
- 2: geneseqp1990s:*
- 3: geneseqp2000s:*
- 4: geneseqp2001s:*
- 5: geneseqp2002s:*
- 6: geneseqp2003as:*
- 7: geneseqp2003bs:*
- 8: geneseqp2004s:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	254	100.0	47	AD061523	AD061523 Human int
2	254	100.0	148	AD061520	AD061520 Human int
3	76	29.9	52	AAB91765	AAB91765 Adrenomed
4	72.5	28.5	52	AAB91759	AAB91759 Adrenomed
5	72.5	28.5	52	AAB75110	AAB75110 Human adr
6	72.5	28.5	52	AAB09818	AAB09818 Human adr
7	72.5	28.5	52	ABP55104	ABP55104 Human adr
8	72.5	28.5	52	ADCC2152	ADCC2152 Human adr
9	72.5	28.5	52	ADG91993	ADG91993 Human mat
10	72.5	28.5	52	ADM94034	ADM94034 Human AM
11	72.5	28.5	52	ADN03278	ADN03278 Exemplary
12	72.5	28.5	52	ADM98309	ADM98309 Mature hu
13	72.5	28.5	52	ADP18429	ADP18429 Neurogene
14	72.5	28.5	52	ADP18402	ADP18402 Neurogene
15	72.5	28.5	52	ADR42113	ADR42113 Adrenomed
16	72.5	28.5	53	AAB75111	AAB75111 Glycine e
17	72.5	28.5	53	AAB75112	AAB75112 Glycine e
18	72.5	28.5	53	ABJ18665	ABJ18665 Universal
19	72.5	28.5	62	AAB75113	AAB75113 Linker hu
20	72.5	28.5	91	ADFS5556	ADFS5556 Active hu
21	72.5	28.5	120	AAB75123	AAB75123 USP4(1-56
22	72.5	28.5	120	AAB75122	AAB75122 USP4(1-57
23	72.5	28.5	120	ABJ18669	ABJ18669 Universal
24	72.5	28.5	120	ABJ18670	ABJ18670 Universal
25	72.5	28.5	147	AAB75124	AAB75124 USP4(1-84

26	72.5	28.5	147	ABJ18671	ABJ18671 Universal
27	72.5	28.5	170	AAB75114	AAB75114 Thiorodox
28	72.5	28.5	185	AAB60344	AAB60344 Human adr
29	72.5	28.5	185	AAB49697	AAB49697 Human adr
30	72.5	28.5	185	ABP72347	ABP72347 Adrenomed
31	72.5	28.5	185	ADA27595	ADA27595 Human adr
32	72.5	28.5	185	ADFS5557	ADFS5557 Human adr
33	72.5	28.5	185	ADN10849	ADN10849 Human adr
34	72.5	28.5	185	ADM98308	ADM98308 Human pre
35	72.5	28.5	185	ADP18855	ADP18855 Human pre
36	72.5	28.5	185	ADP12587	ADP12587 Protein e
37	72.5	28.5	185	ADO36937	ADO36937 Human pro
38	72.5	28.5	185	ADR14147	ADR14147 Human NF-
39	72.5	28.5	185	ADR87608	ADR87608 Human adr
40	72.5	28.5	185	ADP19162	ADP19162 Human adr
41	72.5	28.5	186	ADD18582	ADD18582 Human dis
42	72.5	28.5	206	ABJ18668	ABJ18668 Universal
43	71.5	28.1	184	ABJ57209	ABJ57209 Mouse 18c
44	71.5	28.1	184	ADO60040	ADO60040 CRH signa
45	70.5	27.8	188	AAB60345	AAB60345 Porcine a

ALIGNMENTS

RESULT 1
AD061523 standard; protein; 47 AA.
AD061523;
26-AUG-2004 (first entry)
Human intermedin mature proetin SEQ ID NO:5.
human; intermedin; hypotensive; gene therapy; hypertension;
cardioprotective; diet; prolactin release; growth hormone release;
ovarian follicle survival; oedema.
Homo sapiens.
MO2004048547-A2.
10-JUN-2004.
26-NOV-2003; 2003MO-US037968.
26-NOV-2002; 2002US-0429327P.
(STRD) UNIV LELAND STANFORD JUNIOR.
Hau SYT;
WPI; 2004-44176/41.
New composition comprises an intermedin peptide, useful for treating hypertension, as cardioprotective agent, as a diet aid, for the release of prolactin, or for growth hormone release from the pituitary.
Disclosure; SEQ ID NO 5; 68pp; English.
The invention relates to a novel composition comprising an intermedin peptide, which comprises at least 18 contiguous amino acids in a sequence comprising 148 amino acids (AD061520). A composition of the invention has hypotensive activity, and may have a use in gene therapy. The composition comprising the intermedin peptide is useful for identifying homologous or related genes, for production of the encoded peptide, in producing compositions that modulate the expression or function of its encoded protein, for gene therapy, mapping functional regions of the protein, or in studying associated physiological pathways. The intermedin peptide is useful for treating hypertension, as a cardioprotective agent, as a diet aid, for the release of prolactin, in growth hormone release from the pituitary, or for ovarian follicle survival and growth. They are also

CC useful in the reduction of oedema. The present sequence represents the
CC human intermedin mature protein.

XX
SQ Sequence 47 AA;

Query Match 100.0%; Score 254; DB 8; Length 47;
Best Local Similarity 100.0%; Pred. No. 1.1e-27;
Matches 47; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TQAQLRVGCVLTGTCQVONLSHRLWQMGPAQRQDSAPVDPSSPHSY 47
DB 1 TQAQLRVGCVLTGTCQVONLSHRLWQMGPAQRQDSAPVDPSSPHSY 47

RESULT 2

ADO61520
ID ADO61520 standard; protein; 148 AA.

AC⁻ ADO61520;

XX 26-AUG-2004 (first entry)

DE Human Intermedin SEQ ID NO:2.

XX human; intermedin; hypotensive; gene therapy; hypertension;

KW cardioprotective; diet; prolactin release; growth hormone release;

KM ovarian follicle survival; oedema.

XX Homo sapiens.

PN WO2004048547-A2.

XX 10-JUN-2004.

XX 26-NOV-2003; 2003WO-US037968.

XX 26-NOV-2002; 2002US-0429327P.

XX (STRD) UNIV LELAND STANFORD JUNIOR.

XX Hsu SYT;

XX WPI; 2004-441176/41.

XX N-PSDB; ADO61519.

XX Claim 1; SEQ ID NO 2; 68pp; English.

CC The invention relates to a novel composition comprising an intermedin
CC peptide, which comprises at least 18 contiguous amino acids in a sequence
CC comprising 148 amino acids (ADO61520). A composition of the invention has
CC hypotensive activity, and may have a use in gene therapy. The composition
CC comprising the intermedin peptide is useful for identifying homologous or
CC related genes, for production of the encoded peptide, in producing
CC compositions that modulate the expression or function of its encoded
CC protein, for gene therapy, mapping functional regions of the protein, or
CC in studying associated physiological pathways. The intermedin peptide is
CC useful for treating hypertension, as a cardioprotective agent, as a diet
CC aid, for the release of prolactin, in growth hormone release from the
CC pituitary, or for ovarian follicle survival and growth. They are also
CC useful in the reduction of oedema. The present sequence represents the
CC human intermedin protein of the invention.

XX Sequence 148 AA;

Query Match 100.0%; Score 254; DB 8; Length 148;
Best Local Similarity 100.0%; Pred. No. 3.9e-27;
Matches 47; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TQAQLRVGCVLTGTCQVONLSHRLWQMGPAQRQDSAPVDPSSPHSY 47

DB 101 TQAQLRVGCVLTGTCQVONLSHRLWQMGPAQRQDSAPVDPSSPHSY 147

RESULT 3

AAB91765
ID AAB91765 standard; peptide; 52 AA.

XX AAB91765;

XX 22-JUN-2001 (first entry)

DE Adrenomedullin peptide (AM) SEQ ID NO:941.

KW Protection; endogenous therapeutic peptide; peptidase; conjugation;
KW blood component; modification; succinimideyl; maleimido group; amino;
KW hydroxyl; thiol; hormone; growth factor; neurotransmitter.

XX Homo sapiens.

XX Synthetic.

XX WO200069900-A2.

XX 23-NOV-2000.

XX 17-MAY-2000; 2000WO-US013576.

XX 17-MAY-1999; 99US-0134406P.

XX 10-SEP-1999; 99US-0153406P.

XX 15-OCT-1999; 99US-0159783P.

XX (CONJ-) CONJUCHEM INC.

XX Bridon DP, Ezrin AM, Milner PG, Holmes DL, Thibaudau K;

XX WPI; 2001-112059/12.

XX Modifying and attaching therapeutic peptides to albumin prevents
XX peptidase degradation, useful for increasing length of in vivo activity.

XX Disclosure; Page 501; 733pp; English.

CC The present invention describes a modified therapeutic peptide (I)
CC comprising a therapeutically active amino acid region (III) and a
CC reactive group (II) (e.g. succinimideyl and maleimido groups) attached to
CC a less therapeutically active amino acid region (IV), which covalently
CC bonds with amino/hydroxyl/thiol groups on blood components to form a
CC peptidase stabilised therapeutic peptide composed of 3-50 amino acids.
CC (I) are useful for modifying therapeutic peptides e.g. hormones, growth
CC factors and neurotransmitters, to protect them from peptidase activity in
CC vivo for the treatment of various disorders. Endogenous therapeutic
CC peptides are not suitable as drug candidates as they require frequent
CC administration due to rapid degradation by peptidases in the body.
CC Modifying and attaching therapeutic peptides to albumin prevents or
CC reduces the action of peptidases to increase length of activity (half
CC life) and specificity as bonding to large molecules decreases
CC intracellular uptake and interference with physiological processes.
CC AAB90829 to AAB94441 represent peptides which can be used in the
CC exemplification of the present invention

XX Sequence 52 AA;

Query Match 29.9%; Score 76; DB 4; Length 52;
Best Local Similarity 35.9%; Pred. No. 0.0091;
Matches 14; Conservative 6; Mismatches 19; Indels 0; Gaps 0;

QY 9 GCVLGTQVONLSHRLWQMGPAQRQDSAPVDPSSPHSY 47

DB 14 GCRGTCTVQKLAHQIYQFTDKGVAPRSKRKSKISPGCY 52

RESULT 4

AAB91759

ID	AAB91759 standard; peptide; 52 AA.									
XX										
AC	AAB91759;									
XX										
DT	22-JUN-2001 (first entry)									
XX										
DE	Adrenomedullin peptide (AM) SEQ ID NO:935.									
XX										
KM	Protection; endogenous therapeutic peptide; peptidase; conjugation;									
XX	blood component; modification; succinimidyl; maleimido group; amino;									
KM	hydroxyl; thiol; hormone; growth factor; neurotransmitter.									
XX										
OS	Homo sapiens.									
OS	Synthetic.									
XX										
PN	WO200069900-A2.									
XX										
PD	23-NOV-2000.									
XX										
PF	17-MAY-2000; 2000MO-US013576.									
XX										
PR	17-MAY-1999; 99US-0134406P.									
XX										
PR	10-SEP-1999; 99US-0153406P.									
XX										
PR	15-OCT-1999; 99US-0159783P.									
XX										
PA	(CONU-) CONUCHEM INC.									
XX										
P1	Bridon DP, Ezrin AM, Milner PG, Holmes DL, Thibaudau K;									
XX										
DR	WPI; 2001-112059/12.									
XX										
PT	Modifying and attaching therapeutic peptides to albumin prevents									
PT	peptidase degradation, useful for increasing length of in vivo activity.									
XX										
PS	Disclosure; Page 498; 733pp; English.									
XX										
CC	The present invention describes a modified therapeutic peptide (I)									
CC	comprising a therapeutically active amino acid region (III) and a									
CC	reactive group (II) (e.g. succinimidyl and maleimido groups) attached to									
CC	a less therapeutically active amino acid region (IV), which covalently									
CC	bonds with amino/hydroxyl/thiol groups on blood components to form a									
CC	peptidase stabilised therapeutic peptide composed of 3-50 amino acids.									
CC	(I) are useful for modifying therapeutic peptides e.g. hormones, growth									
CC	factors and neurotransmitters, to protect them from peptidase activity in									
CC	vivo for the treatment of various disorders. Endogenous therapeutic									
CC	peptides are not suitable as drug candidates as they require frequent									
CC	administration due to rapid degradation by peptidases in the body.									
CC	Modifying and attaching therapeutic peptides to albumin prevents or									
CC	reduces the action of peptidases to increase length of activity (half									
CC	life) and specificity as bonding to large molecules decreases									
CC	intracellular uptake and interference with physiological processes.									
CC	AAB90829 to AAB92441 represent peptides which can be used in the									
CC	exemplification of the present invention									
XX										
5Q	Sequence 52 AA;									
XX										
QY	Query Match 28.5%; Score 72.5; DB 4; Length 52;									
DB	Best Local Similarity 38.5%; Pred. No. 0.028; 17; Indels 1; Gaps 1									
XX										
15	GCVLTGTCQVONLSRLMLQMPAGRQSDAPVDPSSPHSY 47									
15	GCRFGCTVQKLAHQIYQPT-DKQDNVAPRKSISPGY 52									
XX										
RESULT 5										
ID	AAB75110									
XX										
AC	AAB75110; standard; protein; 52 AA.									
XX										
DT	31-JUL-2001 (first entry)									
XX										

```
DE Human adrenomedullin (AM) protein.  
XX  
KM Adrenomedullin; glycine extended adrenomedullin; AM; AM-gly;  
KW adrenomedullin precursor; fusion protein; pharmaceutical; diagnostic.  
XX  
OS Homo sapiens.  
XX WO200127310-A1.  
PN 19-APR-2001.  
XX  
PD 10-OCT-2000; 2000WO-JP007023.  
PF 15-OCT-1999; 99JP-00294147.  
PR XX  
PA (SHIO ) SHIONOGI & CO LTD.  
XX  
PI Takimoto A, Mitsuda Y, Nakayama T, Mitsuishima K;  
XX WPI; 2001-282044/29.  
DR N-PADB; AAH19806.  
XX  
PT Producing adrenomedullin useful for pharmaceutical and diagnostic  
PI application comprises producing fused adrenomedullin precursor using a  
recombinant host.  
PX  
PS Disclosure; Page 45; 75pp; Japanese.  
XX  
CC The present invention describes a method (M1) for producing  
adrenomedullin precursor. The method comprises: (a) producing the fused  
protein using a recombinant host cell; (b) restricted digestion of the  
fused protein by a protease followed by collection of sediment; and (c)  
dissolving the sediment and extracting adrenomedullin precursor. The  
method can be used for the production of adrenomedullin precursor for  
pharmaceutical and diagnostic applications. AAH19806 to AAH19866 and  
AAH75110 to AAH75124 represent sequences which are used in the  
embodiment of the present invention  
CX CC  
SC Sequence 52 AA;  
  
Query Match      28.5%; Score 72.5; DB 4; Length 52;  
Best Local Similarity 38.5%; Pred. No. 0.028; Indels 1; Gaps 1;  
Matches 15; Conservative 6; Mismatches 17;  
  
OY          9 GCVTGTCOVONTLSRLWQLMGPPAGRODSAPPVDPSPESHY 47  
DB           |||||:::||::|:  
       15 GCRCGTCTVGKTAHQTYQFT-DKKDKNVAFRSKISPGCY 52  
  
RESULT 6  
AAE09818 ID     AAE09818 standard; peptide: 52 AA.  
XC         AAE09818;  
XD  
XE 29-NOV-2001 (first entry)  
XF  
XG Human adrenomedullin peptide #1.  
XH  
XI  
XJ Homo sap lens.  
XK US6268474-B1.  
XL  
XM 31-Jul-2001.  
XN  
XP 30-Apr-1998;    98US-00070504.  
XQ  
XR 30-Apr-1998;   98US-00070504.  
XS  
XT (UYCR-) UNIT CREIGHTON.
```


RESULT 9
ADG91993
ID ADG91993 standard; protein; 52 AA.
XX
AC ADG91993;
XX
DT 11-MAR-2004 (first entry)
XX
DE Human mature adrenomedullin protein.
XX
KW neuroleptic; antisense therapy; gene therapy; adrenomedullin agonist;
KW schizophrenia; gene expression; decidua protein induced by progesterone;
KW DPP; adrenomedullin; cold shock domain protein A; cda; antisense;
KW siRNA; ribozyme; triple helix formation.
XX
OS Homo sapiens.
XX
PN WO2003078658-A2.
XX
PD 25-SEP-2003.
XX
PF 19-MAR-2003; 2003WO-BP002875.
XX
PR 20-MAR-2002; 2002US-036601P.
XX
PA (NOVS) NOVARTIS AG.
PA (NOVS) NOVARTIS PHARMA GMBH.
XX
PI Buxton FP, Carpenter WT, Roberts RC, Tamminga CA;
XX
DR WPI; 2003-767532/72.
XX
PT Screening for schizophrenia in a population utilizing genes encoding the
PT decidua protein induced by progesterone, adrenomedullin and/or cold
PT shock domain protein A, useful in diagnosing, preventing and/or treating
PT schizophrenia.
XX
XX
PS Disclosure; SEQ ID NO 1; 41pp; English.
XX
CC The invention relates to a method of screening for schizophrenia in a
CC population comprising determining the magnitude of expression, in members
CC of the population, of at least one gene selected from the gene encoding
CC decidua protein induced by progesterone (DPP), the gene encoding
CC adrenomedullin and the gene encoding cold shock domain protein A (cda)
CC in a sample, and comparing the magnitude of expression to a baseline of
CC expression of the gene, where increased gene expression indicates the
CC presence of schizophrenia. An antisense molecule, siRNA, ribozyme or
CC nucleic acid molecule promoting triple helix formation that specifically
CC inhibit the expression of DPP, cda or adrenomedullin gene, is useful
CC for the manufacture of a medicament for the treatment of schizophrenia.
CC An antibody that specifically binds an epitope of DPP, cda or
CC adrenomedullin is also useful for the manufacture of a medicament for the
CC treatment of schizophrenia. This sequence represents the mature
CC adrenomedullin protein.
XX
SQ Sequence 52 AA;
XX
Query Match 28.5%; Score 72.5; DB 7; Length 52;
Best Local Similarity 38.5%; Pred. No. 0.028;
Matches 15; Conservative 6; Mismatches 17; Indels 1; Gaps 1;
OY 9 GCVTGTCOVNLSHRLQWLMGPAGRODAPVDPSPSY 47
DB 15 GCRFGTCTVOKLAHQIYQFT-DKDKXNVAPRSKISPOGY 52

DT 17-JUN-2004 (first entry)
XX
XX Human AM peptide, a CRSP homologue.
DE
XX
KW calcitonin receptor stimulating peptide; CRSP; cAMP activity;
KW skeletal disorder; cancer; hypertension; restenosis; analgesic;
KW appetite suppressant; diuretic; vasotropic; cyostatic; diuretic;
KW osteopathic; anorectic; hypotensive; human; AM.
XX
XX
OS Homo sapiens.
XX
XX
XX Key Location/Qualifiers
FH Disulfide-bond 16..21
FT Modified-site 52
FT /note="C-terminal amide"
XX
XX WO2003102180-A1.
XX
XX
XX 11-DEC-2003.
XX
XX 28-MAY-2003; 2003WO-JP006641.
XX
XX 04-JUN-2002; 2002JP-00162797.
XX
XX (NISC-) JAPAN SCI & TECHNOLOGY CORP.
XX (NINA-) JAPAN NAT CARDIOVASCULAR.
XX
XX Minamino N, Katafuchi T;
XX
XX WPI; 2004-043113/04.
XX
XX
XX Calcitonin receptor stimulating peptide promoting cAMP production in
PT cells for treatment of cancer; skeletal disorders and hypertension and as
PT appetite suppressants and analgesics.
XX
XX
PS Disclosure; Fig 2; 62pp; Japanese.
XX
XX
CC This invention relates to novel calcitonin receptor stimulating peptides
CC (CRSPs) that are expressed in the central nervous system and act on the
CC calcitonin receptor to promote the production of cAMP within a cell.
CC Specifically, it refers to CRSPs that can stimulate concentration
CC dependent sodium ion uptake and furthermore act to inhibit calcium ion
CC uptake by the cell. The present invention describes mutant peptides
CC derived from the CRSPs that have additions, deletions and/or
CC substitutions of one or more amino acids, yet that retain similar
CC activity and are useful for developing drug compositions and
CC pharmaceutically acceptable carriers to treat and/or prevent skeletal
CC disorders, cancer, hypertension and restenosis. Furthermore, CRSPs can
CC also be used as analgesics, appetite suppressants and diuretics such that
CC exhibit vasotropic, cyostatic, diuretic, osteopathic, anorectic and
CC hypotensive activities. This peptide sequence is a human AM peptide that
CC has cAMP activity, given in an exemplification of the invention.
XX
SQ Sequence 52 AA;
XX
Query Match 28.5%; Score 72.5; DB 8; Length 52;
Best Local Similarity 38.5%; Pred. No. 0.028;
Matches 15; Conservative 6; Mismatches 17; Indels 1; Gaps 1;
OY 9 GCVTGTCOVNLSHRLQWLMGPAGRODAPVDPSPSY 47
DB 15 GCRFGTCTVOKLAHQIYQFT-DKDKXNVAPRSKISPOGY 52

RESULT 11
ADN03278
ID ADN03278 standard; peptide; 52 AA.
XX
AC ADN03278;
XX
DT 17-JUN-2004 (first entry)
XX
XX Exemplary peptide ligand for proteome analysis #2.

XX Peptide ligand; proteome; capture compound; mass spectrometry;
 KW protein separation;
 KW matrix assisted laser desorption ionisation-time of flight; MALDI-TOF.
 XX Unidentified.
 OS

PN US2003119021-A1.

XX 26-JUN-2003.

XX 16-JUL-2002; 2002US-00197954.

XX 16-JUL-2001; 2001US-0306019P.

XX 16-AUG-2001; 2001US-0314123P.

XX 11-MAR-2002; 2002US-0363433P.

XX (KOST/) KOSTER H.
 PA (SID/) SIDDIQI S.
 PA (LIT/) LITTLE D P.

XX Koster H, Siddiqi S, Little DP;
 PI

XX WPI; 2004-059185/06.

PT Collection of capture compounds capable of binding to biomolecules to
 PT form complexes that are stable under mass spectrometry conditions, useful
 PT for analysis of biomolecules, especially proteins.

PS Disclosure; SEQ ID NO 2; 165pp; English.

XX The invention relates to a collection of capture compounds capable of
 CC binding to biomolecules to form complexes that are stable under mass
 CC spectrometry conditions. The formulae for the capture compounds comprises
 CC sets of compounds of formula (I)-(III) given in the specification. Also
 CC included are analysis of biomolecules (by contacting a composition
 CC comprising a biomolecule with the above collection and identifying or
 CC detecting bound biomolecules), separating protein conformers (by
 CC collecting a composition comprising a biomolecule with the above
 CC collection, separating the members of the collection and identifying
 CC bound proteins), reducing diversity of a complex mixture of biomolecules
 CC (by contacting the mixture with the above collection and separating each
 CC set of complexes of capture compounds with biomolecules from the other
 CC sets) and identifying phenotype-specific biomolecules (by sorting cells
 CC from a single subject into sets according to a phenotype, contacting
 CC mixtures of biomolecules from each set with the above collection and
 CC comparing the patterns of biomolecule binding from each set). The
 CC collection of capture compounds is useful for the analysis of
 CC biomolecules, especially proteins (e.g. analysis of a proteome), using
 CC mass spectrometry, especially matrix assisted laser desorption ionisation
 CC -time of flight (MALDI-TOF) mass spectrometry. The present sequence is an
 CC exemplary peptide ligand which may be incorporated into a capture
 CC compound of the invention.

XX Sequence 52 AA;

Query Match 28.5%; Score 72.5; DB 8; Length 52;

Best Local Similarity 38.5%; Pred. No. 0.028; Mismatches 17; Indels 1; Gaps 1;

Matches 15; Conservative 6; Mismatches 17; Indels 1; Gaps 1;

DB 9 GCVLTGTCQYQNLSHRLMQLMGPAGRODAPVDPSPHSY 47

15 GCRFGCTCTVQKLAHQIYQFT-DKDXNVAPRSKISPOGY 52

RESULT 12

ID ADM98309 standard; protein; 52 AA.

XX ADM98309;

XX 15-JUL-2004 (first entry)

DE Mature human adrenomedullin protein SeqID 3.

XX AM(11-22); vasoconstriction; human; adrenomedullin; AM; vasoregulatory;

XX blood pressure; vasodilator; vasodilatory shock; septic shock;

XX haemorrhagic shock; vasotropic; hypotensive; immunosuppressive;

XX antibacterial.

OS Homo sapiens.

XX WO2004032708-A2.

XX 22-APR-2004.

XX 03-OCT-2003; 2003WO-US031400.

XX 04-OCT-2002; 2002US-0416291P.

XX (USSH) US DEPT HEALTH & HUMAN SERVICES.

XX Cuttitta F, Martinez A, Stetler-Stevenson WG, Unsworth EJ;

XX Saavedra JM;

XX WPI; 2004-340778/31.

XX New AM(11-22) peptides, useful for inducing vasoconstriction,

XX for treating septic shock, vasodilatory shock or hemorrhagic shock, or

XX for reducing blood pressure.

XX Disclosure; SEQ ID NO 3; 40pp; English.

XX This invention relates to a novel peptide AM(11-22) useful for treating
 CC shock, or in a pharmaceutical composition for inducing vasoconstriction.
 CC Specifically, AM(11-22) is a short peptide derived from human
 CC adrenomedullin (AM), which is a vasoregulatory compound that influences
 CC blood pressure. The present invention describes screening assays to
 CC identify compounds including antibodies, small molecule inhibitors or
 CC peptides that modulate AM(11-22)-mediated vasoconstriction and as such
 CC represent novel vasodilators or vasoconstrictors. Accordingly, AM(11-22)
 CC can be used therapeutically in a pharmaceutical composition to inhibit
 CC blood flow following traumatic or surgical injury, as well as for
 CC vasodilatory, septic or haemorrhagic shock, and thus exhibits vasotropic,
 CC hypotensive, immunosuppressive and antibacterial activities. This
 CC polypeptide sequence is the mature human adrenomedullin protein of the
 CC invention.

XX Sequence 52 AA;

Query Match 28.5%; Score 72.5; DB 8; Length 52;

Best Local Similarity 38.5%; Pred. No. 0.028; Mismatches 17; Indels 1; Gaps 1;

Matches 15; Conservative 6; Mismatches 17; Indels 1; Gaps 1;

DB 9 GCVLTGTCQYQNLSHRLMQLMGPAGRODAPVDPSPHSY 47

15 GCRFGCTCTVQKLAHQIYQFT-DKDXNVAPRSKISPOGY 52

RESULT 13

ID ADP18429 standard; protein; 52 AA.

XX ADP18429;

XX 26-AUG-2004 (first entry)

DE Neurogenesis modulation-related protein SeqID43.

XX neurogenesis modulation; neural tissue; central nervous system disorder;

XX neurodegenerative; ischemic; learning and memory disorder;

XX neurological trauma; nootropic; neuroprotective; CNS-gen;

XX cerebrotropic; vasotropic; anticonvulsant; antiparkinsonian;

XX haemostatic; hypertensive; muscular-gen; ophthalmological;

XX antiinflammatory; analgesic; antidiabetic; neurogenesis modulator;

XX neural stem cell; progenitor cell;

neural tissue G-protein coupled receptor activator; neurogenesis inducer; intracellular neural Ca 2+ enhancer; intracellular neural cAMP stimulator; intracellular neural Ca 2+ enhancer; Parkinson's disease; Parkinson's disorder; Huntington's disease; Alzheimer's disease; multiple sclerosis; amyotrophic lateral sclerosis; Shy-Drager syndrome; progressive supranuclear palsy; Lewy body disease; spinal ischaemia; ischaemic stroke; cerebral infarction; spinal cord injury; cancer-related brain; spinal cord injury; multi-infarct dementia; geriatric dementia; cAMP level; embryonic tissue; human.

Homo sapiens.

MO2004045592-A2.

03-JUN-2004.

20-NOV-2003; 2003MO-IB005311.

20-NOV-2002; 2002US-0427912P.

(NEUR-) NEURONOVA AB.

(BERT) BERTILSSON G.

(ERLA) ERLANDSSON R.

(FRIS) FRISSEN J.

(HAEG) HAEGSTRAND A.

(HEID) HEIDRICH J.

(HEIL) HEILSTROM K.

(HAEG) HAEGBLAD J.

(JANS) JANSSEN K.

(KORT) KORTESMAA J.

(LUND) LINDQUIST P.

(LUND) LUNDH H.

(MCGU) MCGUIRE J.

(MERC) MERCER A.

(NJBG) NJBERG K.

(OSSO) OSSOINAK A.

(PATR) PATRONE C.

(ROEN) ROENHOLM H.

(ZACH) ZACHRISSON O.

(WIKS) WIKSTROM L.

Bertilsson G, Eriandsson R, Frisen J, Haegestranda A, Heidrich J, Hellstroem K, Haegblad J, Jansson K, Kortesmaa J, Lindquist P, Lundh H, Meguire J, Mercer A, Njberg K, Ossoinak A, Patrone C, Roenholm H, Zachrisson O, Wikstrom L;

WPI; 2004-449666/42.

Use of agent(s) that elevate intracellular cyclic adenosine monophosphate or intracellular calcium levels in neural tissue for modulating neurogenesis to treat central nervous system disorder.

Disclosure; SEQ ID NO 43; 77pp; English.

This invention relates to a novel method of modulating neurogenesis in the neural tissue of a patient exhibiting symptom(s) of a central nervous system disorder, such as neurodegenerative, ischaemic or learning and memory disorder or neurological trauma. The method involves at least one agent (A) that elevates intracellular cyclic adenosine monophosphate (cAMP) levels or at least one agent (B) that elevates intracellular Ca 2+ levels in the neural tissue, which is administered where (A) modulates and (B) induces neurogenesis. The invention may be useful for the production of compounds with a neurotropic, neuroprotective, CNS-Gen, cerebroprotective, vasotropic, anticonvulsant, antiparkinsonian, haemostatic, hypertensive, muscular-Gen, ophthalmological, antiinflammatory, analgesic or antidiabetic activity. These compounds may act as neurogenesis modulators, neural stem or progenitor cell proliferation, differentiation and/or migration modulators, neural tissue G-protein coupled receptor activators, neurogenesis inducers, intracellular neural cAMP enhancers, intracellular neural Ca 2+ stimulators or intracellular Ca 2+ enhancers. The invention is useful for modulating neurogenesis in neural tissue of a patient exhibiting at least one symptom of central nervous system disorder, such

as Parkinson's disease and Parkinson's disorders, Huntington's disease, Alzheimer's disease, multiple sclerosis, amyotrophic lateral sclerosis, Shy-Drager syndrome, progressive supranuclear palsy, Lewy body disease, spinal ischaemia, ischaemic stroke, cerebral infarction, spinal cord injury, cancer-related brain and spinal cord injury, multi-infarct dementia, and geriatric dementia; for increasing the intracellular levels of or stimulating cAMP levels in a cell (preferably a cell from a neural tissue); and for in vitro modulation of neurogenesis. The agent modulates neurogenesis in neural tissue by modulating proliferation, differentiation, migration or survival of neural stem cells or progenitor cells in the tissue; by maintaining or increasing the amount or percentage of doublecortin positive cells in the neural tissue relative to a patient not dosed with the agent or by activation of a G-protein coupled receptor in the neural tissue. The method results in elevation of cAMP levels of the neural stem cells by over 20% compared to untreated tissue. The in vivo induction of neurogenesis allows treatment of disorders caused by cell loss, injury or disease by endogenous replacement and obviates the need for transplanting foreign cells into a patient. Neurogenesis can also be induced by administration of the neurogenesis-modulating agent directly into a desired site, which avoids unnecessary systemic administration and possible side effects and further provides an alternative to the use of drugs and the controversial use of large quantities of embryonic tissue for treatment of Parkinson's disease. The present sequence is that of a polypeptide which has been shown to have the ability to increase intracellular cAMP levels and which is related to the method of the invention.

Sequence 52 AA;

Query Match 28.5%; Score 72.5; DB 8; Length 52;

Best Local Similarity 38.5%; Pred. No. 0.028;

Matches 15; Conservative 6; Mismatches 17; Indels 1; Gaps 1;

Qy 9 GCVLTGCVQVNLHRLMQLMGPRGDSAPVDSSPSHY 47

Db 15 GCRFGTCVQKLAHQIVQFT-DXDKVAVPRSKISPOGY 52

RESULT 14

ID ADP18402 standard; protein; 52 AA.

AC ADP18402;

DT 26-AUG-2004 (first entry)

XX Neurogenesis modulation-related protein Segidp16.

XX neurogenesis modulation; neural tissue; central nervous system disorder;

KM neurodegenerative; ischaemic; learning and memory disorder;

KM neurological trauma; neurotropic; neuroprotective; CNS-Gen;

KM cerebroprotective; vasotropic; anticonvulsant; antiparkinsonian;

KM haemostatic; hypertensive; muscular-Gen; ophthalmological;

KM antiinflammatory; analgesic; antidiabetic; neurogenesis modulator;

KM neural stem cell; progenitor cell;

KM neural tissue G-protein coupled receptor activator; neurogenesis inducer;

KM intracellular neural cAMP enhancer; intracellular neural cAMP stimulator;

KM intracellular neural Ca 2+ enhancer; Parkinson's disease;

KM Parkinson's disorder; Huntington's disease; Alzheimer's disease;

KM multiple sclerosis; amyotrophic lateral sclerosis; Shy-Drager syndrome;

KM progressive supranuclear palsy; Lewy body disease; spinal ischaemia;

KM ischaemic stroke; cerebral infarction; spinal cord injury;

KM cancer-related brain; spinal cord injury; multi-infarct dementia;

KM geriatric dementia; cAMP level; embryonic tissue; human.

OS Homo sapiens.

XX

PN MO2004045592-A2.

XX

PD 03-JUN-2004.

XX

PF 20-NOV-2003; 2003MO-IB005311.

XX

PR 20-NOV-2002; 2002US-0427912P.

XX (NEUR-) NEURONOVA AB.
 PA (BERT) BERTILSSON G.
 PA (ERLA) ERLANDSSON R.
 PA (FRIS) FRISSEN J.
 PA (HAEG) HAEGSTRAND A.
 PA (HEID) HEIDRICH J.
 PA (HELL) HELSTROEM K.
 PA (HAEG) HAEGBLAD J.
 PA (JANS) JANSSEN K.
 PA (KORT) KORTESMAA J.
 PA (LIND) LINDQUIST P.
 PA (LUND) LUNDH H.
 PA (MCGU) MCGUIRE J.
 PA (MERC) MERCER A.
 PA (NUBE) NUBERG K.
 PA (OSSO) OSSOINAK A.
 PA (PATR) PATRONE C.
 PA (ROEN) ROENNHOLOM H.
 PA (ZACH) ZACHRISSON O.
 PA (WIKS) WIKSTROEM L.

XX Bertilsson G, Erlandsson R, Frisen J, Haegstrand A, Heidrich J;
 PI Hellstroem K, Haegblad J, Jansson K, Kortesmaa J, Lindquist P;
 PI Lundh H, McGuire J, Mercer A, Njberg K, Ossoinak A, Patrone C;
 PI Roennholm H, Zachrisson O, Wikstroem L;
 DR WPI; 2004-449666/42.

PT Use of agent (s) that elevate intracellular cyclic adenosine monophosphate
 or intracellular calcium levels in neural tissue for modulating
 PT neurogenesis to treat central nervous system disorder.

PS Disclosure; SEQ ID NO 16; 77bp; English.

XX This invention relates to a novel method of modulating neurogenesis in
 CC the neural tissue of a patient exhibiting symptom(s) of a central nervous
 CC system disorder, such as neurodegenerative, ischemic or learning and
 CC memory disorder or neurological trauma. The method involves at least one
 CC agent (A) that elevates intracellular cyclic adenosine monophosphate
 CC (cAMP) levels or at least one agent (B) that elevates intracellular Ca 2+
 CC levels in the neural tissue, which is administered where (A) modulates
 CC and (B) induces neurogenesis. The invention may be useful for the
 CC production of compounds with a nootropic, neuroprotective, CNS-Gen,
 CC cerebroprotective, vasotropic, anticonvulsant, antiparkinsonian,
 CC haemostatic, hypertensive, muscular-Gen, ophthalmological,
 CC antiinflammatory, analgesic or antidiabetic activity. These compounds may
 CC act as neurogenesis modulators, neural stem or progenitor cell
 CC proliferation, differentiation and/or migration modulators, neural tissue
 CC G-protein coupled receptor activators, neurogenesis inducers,
 CC intracellular neural cAMP enhancers, intracellular neural cAMP
 CC stimulators or intracellular neural Ca 2+ enhancers. The invention is
 CC useful for modulating neurogenesis in neural tissue of a patient
 CC exhibiting at least one symptom of central nervous system disorder, such
 CC as Parkinson's disease and Parkinson's disorders, Huntington's disease,
 CC Alzheimer's disease, multiple sclerosis, amyotrophic lateral sclerosis,
 CC Shy-Drager syndrome, progressive supranuclear palsy, Lewy body disease,
 CC spinal ischaemia, ischaemic stroke, cerebral infarction, spinal cord
 CC injury, cancer-related brain and spinal cord injury, multi-infarct
 CC dementia and geriatric dementia; for increasing the intracellular levels
 CC of or stimulating cAMP levels in a cell (preferably a cell from a neural
 CC tissue); and for in vitro modulation of neurogenesis. The agent modulates
 CC neurogenesis in neural tissue by modulating proliferation,
 CC differentiation, migration or survival of neural stem cells or progenitor
 CC cells in the tissue; by maintaining or increasing the amount or
 CC percentage of doublecortin positive cells in the neural tissue relative
 CC to a patient not doublecortin positive cells in the neural tissue of a G-protein
 CC coupled receptor in the neural tissue. The method results in elevation of
 CC cAMP levels of the neural stem cells by over 20% compared to untreated
 CC tissue. The in vivo induction of neurogenesis allows treatment of
 CC disorders caused by cell loss, injury or disease by endogenous
 CC replacement and obviates the need for transplanting foreign cells into a

CC patient. Neurogenesis can also be induced by administration of the
 CC neurogenesis-modulating agent directly into a desired site, which avoids
 CC unnecessary systemic administration and possible side effects and further
 CC provides an alternative to the use of drugs and the controversial use of
 CC large quantities of embryonic tissue for treatment of Parkinson's
 CC disease. The present sequence is that of a polypeptide which has been
 CC shown to have the ability to increase intracellular cAMP levels and which
 CC is related to the method of the invention.

XX Sequence 52 AA;

Query Match 28.5%; Score 72.5; DB 8; Length 52;

Best Local Similarity 38.5%; Pred. No. 0.028; Mismatches 15; Conservative 6; Mismatches 17; Indels 1; Gaps 1;

QY 9 GCVLGTCQYQVNTSHRLMQLMGPAQRQDSAPVDPSPHSY 47
 DB 15 GCRFGCTVQKLAHQIYQFT-DKDXDNVAPRSKISPOGY 52

RESULT 15

ID ADR42113 standard; peptide; 52 AA.

AC ADR42113;

DT 21-OCT-2004 (first entry)

DE Adrenomedullin related peptide ligand, SEQ ID 2.

KW Human; ligand; Adrenomedullin.

OS Homo sapiens.

PN WO2004064972-A2.

PD 05-AUG-2004.

PF 16-JAN-2004; 2004WO-US0001037.

PR 16-JAN-2003; 2003US-0441398P.

PA (HKPH-) HK PHARM INC.

PA (KOSK-) KOESTER H.

PI Koester H, Little DP, Siddiqi SM, Greallish MP, Marappan S;

PI Haseman CF, Yip P;

DR WPI; 2004-642213/62.

PT Identifying drug non-target biomolecules in mixture of biomolecules
 PT involves interacting mixture of biomolecules with capture compounds
 PT having high binding affinity and analyzing captured biomolecules to
 PT identify drug non-targets.

PS Disclosure; SEQ ID NO 2; 368bp; English.

XX The present invention relates to a method for identifying drug non-target
 CC biomolecules in a mixture of biomolecules. The method comprises
 CC interacting mixture with capture compounds having moiety X which
 CC covalently binds to biomolecules with high affinity, moiety Y that
 CC increases selectivity of binding so that the capture compound binds to
 CC fewer biomolecules, and moiety Z for presenting X and Y, and analyzing
 CC captured biomolecules to identify drug non-targets. The capture compound
 CC also optionally comprises a sorting function moiety Q and or a solubility
 CC function moiety W. The selectivity function moiety Y serves to modulate
 CC the reactivity function by reducing the number of groups to which the
 CC reactivity function moiety X binds, such as by steric hindrance and other
 CC interactions. Y is optionally a peptide ligand (ADR42112-ADR42256).

XX Sequence 52 AA;

Query Match 28.5%; Score 72.5; DB 8; Length 52;

Best Local Similarity 38.5%; Pred. No. 0.028;
Matches 15; Conservative 6; Mismatches 17; Indels 1; Gaps 1;

QY 9 GCCTGTCQVONTLSHRLMQLMGPAQRDSAPVDPSSPHSY 47
Db 15 GCRFGCTVOKLAHQIYQFT-DKQDNVAPRSKISPOGY 52

Search completed: May 4, 2005, 18:11:09
Job time : 164 secs

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GenCore version 5.1.6
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OM protein - protein search, using sw model

Run on: May 4, 2005, 17:57:34 ; Search time 176 Seconds
(without alignments)
136.748 Million cell updates/sec

Title: US-10-723-368-5

Perfect score: 254

Sequence: 1 TQQLRVGCVLGTQVQNL.....MGPRGRDAPVDPSSPHSY 47

Scoring table: BLOSUM62

Gapop 10.0 , Gapext 0.5

Searched: 1612378 seqs, 512079187 residues

Total number of hits satisfying chosen parameters: 1612378

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database: UniProt 03:*

1: uniprot_sprot:*

2: uniprot_trembl:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	254	100.0	148	1	ADM2_HUMAN
2	228	89.8	150	1	ADM2_MOUSE
3	225	88.6	146	1	ADM2_RAT
4	188	74.0	159	2	Q75XW6
5	187	73.6	168	2	Q75XW7
6	185.5	73.0	140	2	Q61F59
7	87.5	34.4	174	2	Q75XW8
8	83	32.7	123	2	Q75XW4
9	78.5	30.9	171	2	Q61R55
10	72.5	28.5	185	1	ADM1_HUMAN
11	71.5	28.1	184	1	ADM1_MOUSE
12	71.5	28.1	188	1	ADM1_BOVIN
13	71.5	28.1	188	2	Q95K00
14	70.5	27.8	188	1	ADM1_PIG
15	70	27.6	927	2	Q75S75
16	69.5	27.4	185	1	ADM1_RAT
17	69.5	27.4	188	1	ADM1_CANFA
18	64	25.2	388	2	Q68Y70
19	63	24.8	798	2	Q68Y70
20	59	23.2	142	2	Q6P390
21	59	23.2	386	2	Q7ZXW1
22	59	23.2	506	2	Q6D6B0
23	58.5	23.0	576	2	Q8N2F7
24	58.5	23.0	1281	2	Q6NRJ9
25	58	22.8	281	2	Q93RX1
26	58	22.8	339	2	Q8YCG6
27	58	22.8	351	2	Q6TRC2
28	58	22.8	516	2	Q9P775
29	58	22.8	551	2	Q76912
30	58	22.8	557	2	Q8G1J5
31	58	22.8	559	2	Q9QW71

32	58	22.8	1306	2	Q95QM6	Q95QM6 caenorhabdi
33	58	22.8	1544	2	Q19194	Q19194 caenorhabdi
34	58	22.8	1557	2	Q8WPK9	Q8WPK9 oikopleura
35	57	22.4	82	2	Q6ERN3	Q6ERN3 cryza saciv
36	57	22.4	810	1	ZCC5_HUMAN	Q9C0U0 homo sapien
37	57	22.4	810	1	ZCC5_MOUSE	Q9C0U0 mus sapien
38	57	22.4	810	2	Q6E222	Q6E222 canis famli
39	57	22.4	811	2	Q7ZWS6	Q7ZWS6 xenopus lae
40	57	22.4	1057	2	Q70129	Q70129 anopheles g
41	57	22.4	1784	2	Q94606	Q94606 leishmania
42	56.5	22.2	521	2	Q61NA0	Q61NA0 homo sapien
43	56	22.0	184	2	Q92N11	Q92N11 rhizobium m
44	55.5	21.9	411	2	Q99PY2	Q99PY2 streptomyce
45	55.5	21.9	528	2	Q8BRJ4	Q8BRJ4 pseudomonas

ALIGNMENTS

RESULT 1	ADM2_HUMAN	STANDARD	PRT	148 AA.
ID	ADM2_HUMAN	Q7Z4H4		
AC	Q7Z4H4			
DT	05-JUL-2004 (Rel. 44, Created)			
DT	05-JUL-2004 (Rel. 44, Last sequence update)			
DT	25-OCT-2004 (Rel. 45, Last annotation update)			
DE	Adrenomedullin 2 precursor (intermedin) [contains: Adrenomedullin 2 (intermedin-long) (IMDL), intermedin-short (IMDS)].			
DE	(Intermedin-long) (IMDL); intermedin-short (IMDS)].			
GN	Homo sapiens (Human).			
OS	Homo sapiens (Human).			
OC	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;			
OC	Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.			
OX	NCBI_TaxID=9606;			
RN	[1]			
RP	SEQUENCE FROM N.A., FUNCTION, AND TISSUE SPECIFICITY.			
RC	PubMed=1615490; DOI=10.1074/jbc.M30532200;			
RX	PubMed=14706825; DOI=10.1016/S0014-5793(03)01368-1;			
RA	Takel Y., Inoue K., Ogoshi M., Kawahara T., Bannai H., Miyano S.;			
RT	"Identification of novel adrenomedullin in mammals: a potent cardiovascular and renal regulator.";			
RT	cardiovascular and renal regulator.";			
RL	FEBS Lett. 556:53-58(2004).			
RN	[3]			
RP	SEQUENCE FROM N.A.			
RX	MDLINE=20057165; PubMed=10591208; DOI=10.1038/990031;			
RA	Dunham I., Hunt A.R., Collins J.E., Bruskewich R., Beare D.M.,			
RA	Camp M., Smith L.J., Alnecough R., Almeida J.P., Babbage A.K.,			
RA	Bagnoley C., Bailey J., Barlow K.F., Bates K.N., Beasley O.P.,			
RA	Bird C.P., Blakey S.E., Bridgman A.M., Buck D., Burgess J., Clark G.,			
RA	Burill W.D., Burton J., Carder C., Carter N.P., Chen Y., Clark G.,			
RA	Clegg S.M., Cobley V.E., Cole C.G., Collier R.E., Connor R.,			
RA	Conroy D., Corby N.R., Coville G.J., Cox A.V., Davis J., Dawson E.,			
RA	Dami P.D., Dockree C., Dodsworth S.J., Durbin R.M., Ellington A.G.,			
RA	Evans K.L., Fey J.M., Fleming K., French L., Garner A.A.,			
RA	Gilbert J.G.R., Goward M.E., Graham D.V., Griffiths M.N.D., Hall C.,			
RA	Hall R.E., Hall-Tamlyn G., Heathcote R.W., Ho S., Holmes S.,			
RA	Hunt S.E., Jones M.C., Kershaw J., Kimberley A.M., King A.,			
RA	Laird G.K., Langford C.F., Leversha M.A., Lloyd C., Lloyd D.M.,			
RA	Martyn I.D., Mashreghi-Mohammadi M., Matthews L.H., McCann O.T.,			
RA	McClay L., McLaren S., McMurray A.A., Milne S.A., Mortimore B.J.,			
RA	McClay C.N., Pavitt R., Pearce A.V., Pearson D., Phillimore B.J.C.T.,			
RA	Phillips S.H., Plumb R.W., Ramsey H., Ramsey Y., Rogers L., Ross M.T.,			
RA	Scott C.E., Sena H.K., Skuce C.D., Smalley S., Smith W.L.,			
RA	Soderlund C., Spragon L., Stewart C.A., Sultson J.E., Swann R.M.,			
RA	Vaudin M., Wall M., Wallis J.M., Whiteley M.N., Willey D.L.,			
RA	Williams L., Williams S.A., Williamson H., Wilmer T.E., Wilming L.,			

RA Wright C.L., Hubbard T., Bentley D.R., Beck S., Rogers J., Shimizu N.,
RA Minoshima S., Kawasaki K., Sasaki T., Asakawa S., Kodoh J.,
RA Shintani A., Shibuya K., Yoshizaki Y., Aoki N., Mitsuyama S.,
RA Roe B.A., Chen F., Chu L., Crabtree J., Deschamps S., Do A., Do T.,
RA Dorman A., Fang F., Fu Y., Hua A., Kenton S., Lai H., Lao H.I.,
RA Lewis J., Lewis S., Lin S.-P., Loh P., Malaj E., Nguyen T., Pan H.,
RA Phan S., Qi S., Qian Y., Ray L., Ren Q., Shalli S., Sloan D., Song L.,
RA Wang Q., Wang Y., Wang Z., White J., Willingham D., Wu H., Yao Z.,
RA Zhan M., Zhang G., Chisoe S., Murray J., Miller N., Mink P.,
RA Fulton R., Johnson D., Bemis G., Bentley D., Bradshaw H., Bourne S.,
RA Cordes M., Du Z., Fulton L., Goela D., Graves T., Hawkins J.,
RA Hinds K., Kemp K., Latreille P., Layman D., Ozersky P., Rohlfing T.,
RA Scheet P., Walker C., Wamsley A., Wohldmann P., Pepin K., Nelson J.,
RA Korf I., Bedell J.A., Hallier L.W., Mardis E., Waterston R.,
RA Wilson R., Emanuel B.S., Shaikh T., Kuraishi H., Saita S.,
RA Budarf M.L., McDermid H.E., Johnson A., Wong A.C.C., Morrow B.E.,
RA Edelmann L., Kim U.J., Saitzuya H., Simon M.I., Dumanek J.P.,
RA O'Brien K.P., Wilkinson P., Bodenteich A., Tapia I., Bruder C.E.,
RA Khan A.S., Lane L., Tilaun Y., Wright H., Hartman K., Hu X.,
RA "The DNA sequence of human chromosome 22.";
RT Nature 402:489-495(1999).
CC -1- FUNCTION: IMDL and IMDS may play a role as physiological
CC regulators of gastrointestinal, cardiovascular bioactivities
CC mediated by the CALCRL/RAMPs receptor complexes. Activates the
CC CAMP-dependent pathway.
CC -1- SUBCELLULAR LOCATION: Secreted.
CC -1- TISSUE SPECIFICITY: Expressed in the esophagus, stomach, jejunum,
CC ileum, ileocecum, ascending colon, transverse colon, descending
CC colon and rectum.
CC -1- SIMILARITY: Belongs to the adrenomedullin family.
CC -----
CC This SWISS-PROT entry is copyright. It is produced through a collaboration
CC between the Swiss Institute of Bioinformatics and the EMBL outstation
CC the European Bioinformatics Institute. There are no restrictions on its
CC use by non-profit institutions as long as its content is in no way
CC modified and this statement is not removed. Usage by and for commercial
CC entities requires a license agreement (See <http://www.isb-sib.ch/announce/>
CC or send an email to license@isb-sib.ch).
CC -----
CC EMBL: AF529213; AAC09100.1; -;
CC EMBL: AB121034; BAD07411.1; -;
CC EMBL: AL096767; -; NOT ANNOTATED_CDS.
CC Genew; HGNC:28898; ADM2.
CC MIM: 608682; -;
CC DR Amdaction; Cleavage on pair of basic residues; Hormone; Signal.
CC KW SIGNAL: 1 24
CC FT PROPEP 25 98
CC FT PEPTIDE 101 147 By similarity.
CC FT PEPTIDE 108 147 Adrenomedullin 2 (By similarity).
CC FT DISULFID 110 115 Intermedin-short (Potential).
CC FT MOD_RES 147 147 Tyrosine amide (G-148 provides amide
CC group) (Probable).
CC SQ SEQUENCE 148 AA; 15865 MW; 6E0E3098CCE5BF2 CRC64;
CC
CC Query Match 100.0%; Score 254; DB 1; Length 148;
CC Best Local Similarity 100.0%; Pred. No. 1.2e-24;
CC Matches 47; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
CC
CC 1 TQAQLRVCVIGTCOVONTSHRLWQMGPRGRDSAPVDPSSPSHY 47
CC 101 TQAQLRVCVIGTCOVONTSHRLWQMGPRGRDSAPVDPSSPSHY 147
CC
CC RESULT 2
CC ADM2_MOUSE
CC ID ADM2_MOUSE STANDARD; PRT; 150 AA.
CC AC Q7TNK8;
CC DT 05-JUL-2004 (Rel. 44, Created)
CC DT 05-JUL-2004 (Rel. 44, Last sequence update)
CC DT 05-JUL-2004 (Rel. 44, Last annotation update)
CC DE Adrenomedullin 2 precursor (Intermedin) (Contains: Adrenomedullin 2
CC (Intermedin-long) (IMDL); Intermedin-short (IMDS)).

CN Name=Adm2; Synonym=Am2;
OS Mus musculus (Mouse).
OC Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;
OC Mammalia; Euteleostomi; Rodentia; Sciurognathi; Muridae; Mus.
OX NCBI_Taxid=10090;
CX (1)
RP SEQUENCE FROM N.A., FUNCTION, AND TISSUE SPECIFICITY.
RC STRAIN=C57BL/6;
RC PubMed=14615490; DOI=10.1074/jbc.M305332200;
RT Roh J., Chang C.L., Bhalla A., Klein C., Hau S.Y.T.,
RT "Intermedin is a calcitonin/calcitonin gene-related peptide family
RT peptide acting through the calcitonin receptor-like receptor/receptor
RT activity-modifying protein receptor complexes.";
RT J. Biol. Chem. 279:7264-7274(2004).
RN [2]
RP SEQUENCE FROM N.A., FUNCTION, AND TISSUE SPECIFICITY.
RC TISSUE=Kidney;
RX PubMed=14706825; DOI=10.1016/S0014-5793(03)01368-1;
RA Taketani Y., Inoue K., Ogoshi M., Kawahara T., Bannai H., Miyano S.,
RT "Identification of novel adrenomedullin in mammals: a potent
RT cardiovascular and renal regulator.";
RT FEBS Lett. 536:53-58(2004).
CC -1- FUNCTION: IMDL and IMDS may play a role as physiological
CC regulators of gastrointestinal, cardiovascular bioactivities
CC mediated by the CALCRL/RAMPs receptor complexes. Activates the
CC CAMP-dependent pathway.
CC -1- SUBCELLULAR LOCATION: Secreted.
CC -1- TISSUE SPECIFICITY: High expression detected in the submaxillary
CC gland, kidney, stomach, and mesentery, followed by the pituitary,
CC lung, pancreas, intestine, spleen, thymus and ovary. Expressed
CC mainly in the intermediate lobe of the pituitary, with sporadic in
CC the anterior lobe.
CC -1- SIMILARITY: Belongs to the adrenomedullin family.
CC -----
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CC or send an email to license@isb-sib.ch).
CC -----
CC EMBL: AF529213; AAC09099.1; -;
CC EMBL: AB121035; BAD07412.1; -;
CC MGD: MGI:2675256; Adm2.
CC DR Amdaction; Cleavage on pair of basic residues; Hormone; Signal.
CC KW SIGNAL: 1 25
CC FT PROPEP 26 100
CC FT PEPTIDE 103 149 By similarity.
CC FT PEPTIDE 110 149 Adrenomedullin 2 (By similarity).
CC FT DISULFID 112 117 Intermedin-short (Potential).
CC FT MOD_RES 149 149 Tyrosine amide (G-150 provides amide
CC group) (Probable).
CC SQ SEQUENCE 150 AA; 16188 MW; 2BF3928BFDEBBA CRC64;
CC
CC Query Match 89.8%; Score 228; DB 1; Length 150;
CC Best Local Similarity 93.3%; Pred. No. 2.6e-21;
CC Matches 42; Conservative 2; Mismatches 1; Indels 0; Gaps 0;
CC
CC 3 AQLIRVCVIGTCOVONTSHRLWQMGPRGRDSAPVDPSSPSHY 47
CC 105 AQLIRVCVIGTCOVONTSHRLWQMGPRGRDSAPVDPSSPSHY 149
CC
CC RESULT 3
CC ADM2_RAT
CC ID ADM2_RAT STANDARD; PRT; 146 AA.
CC AC P61312;
CC DT 05-JUL-2004 (Rel. 44, Created)
CC DT 05-JUL-2004 (Rel. 44, Last sequence update)
CC DT 05-JUL-2004 (Rel. 44, Last annotation update)
CC DE Adrenomedullin 2 precursor (Intermedin) (Contains: Adrenomedullin 2
CC (Intermedin-long) (IMDL); Intermedin-short (IMDS)).

GN Name=Adm2; Synonym=Am2;
OS Ratus norvegicus (Rat).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Rattus.
RN NCBI_TaxID=10116;
RP SEQUENCE FROM N.A.
RC TISSUE=Kidney;
RX PubMed=14706825; DOI=10.1016/S0014-5793(03)01368-1;
RA Takei Y., Inoue K., Ogoishi M., Kawahara T., Bannai H., Miyano S.;
RT Identification of novel adrenomedullin in mammals: a potent
RL Cardiovascular and renal regulator.";
FEBS Lett. 556:53-58(2004).
RN
RP FUNCTION, AND TISSUE SPECIFICITY.
RX PubMed=14615490; DOI=10.1074/jbc.M305332200;
RA Roh J., Chang C.L., Bhalla A., Klein C., Hsu S.Y.T.;
RT Intermedin is a calcitonin/calcitonin gene-related peptide family
RT peptide acting through the calcitonin receptor-like receptor/receptor
RT activity-modifying protein receptor complexes.";
J. Biol. Chem. 279:7264-7274(2004).
RL
CC -1- FUNCTION: IMDL and IMDS may play a role as physiological
CC regulators of gastrointestinal, cardiovascular bioactivities
CC mediated by the CALCR/RAMPs receptor complexes. Activates the
CC CAMP-dependent pathway.
CC -1- SUBCELLULAR LOCATION: Secreted.
CC -1- TISSUE SPECIFICITY: Expression was restricted to the intermediate
CC and anterior lobes of the pituitary.
CC -1- SIMILARITY: Belongs to the adrenomedullin family.
CC
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CC or send an email to license@ebi.ac.uk).

DR EMBL: AB121036; BAD07413.1; -
KW Amidation; Cleavage on pair of basic residues; Hormone; Signal.
FT SIGNAL 1 25 Potential.
FT PROPEP 26 96 By similarity.
FT PEPTIDE 99 145 Adrenomedullin 2 (By similarity).
FT PEPTIDE 106 145 Intermedin-short (Potential).
FT DISULFID 108 113 By similarity.
FT MOD_RES 145 145 Tyrosine amide (G-146 provides amide
group) (Probable).
SQ SEQUENCE 146 AA; 15572 MW; C87043237AD29DDC CRC64;

Query Match 88.6%; Score 225; DB 1; Length 146;
Best Local Similarity 91.1%; Pred. No. 6.2e-21;
Matches 41; Conservative 3; Mismatches 1; Indels 0; Gaps 0;

QY 3 AQLRVCVLTGTCOVONLSHRLMQLMGAPGRDSPA PVPSPSPHSY 47
DB 101 AQLRVCVLTGTCOVONLSHRLMQLVAPSGRDSAPVPSPSPHSY 145

RESULT 4
Q75XW6 PRELIMINARY; PRT; 159 AA.
AC Q75XW6;
DT 05-JUL-2004 (TEMBLrel. 27, Created)
DT 05-JUL-2004 (TEMBLrel. 27, Last sequence update)
DE 05-JUL-2004 (TEMBLrel. 27, Last annotation update)
DB Adrenomedullin-3.
GN Name=ADM2;
OS Rugu rubripes (Japanese pufferfish) (Takifugu rubripes).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Actinopterygii; Neopterygii; Teleostei; Euteleostei; Neoteleostei;
OC Acanthomorphia; Acanthopterygii; Percomorpha; Tetraodontiformes;
OC Tetraodontidae; Tetraodontidae; Takifugu.
RN NCBI_TaxID=31033;

RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=22984567; PubMed=14623291; DOI=10.1016/j.bbrcc.2003.10.111;
RA Ogoishi M., Inoue K., Takei Y.,
RT Identification of a novel adrenomedullin gene family in teleost
RT fish.";
RL Biochem. Biophys. Res. Commun. 311:1072-1077(2003).
DR EMBL: AB120297; BAD02343.1; -
SQ SEQUENCE 159 AA; 18028 MW; F1863B20100E254D CRC64;

QY 4 QLRVCVLTGTCOVONLSHRLMQLMGAPGRDSPA PVPSPSPHSY 47
DB 115 QLRVCVLTGTCOVONLSHRLMQLMGAPGRDSPA PVPSPSPHSY 158

RESULT 5
Q75XW7 PRELIMINARY; PRT; 168 AA.
AC Q75XW7;
DT 05-JUL-2004 (TEMBLrel. 27, Created)
DT 05-JUL-2004 (TEMBLrel. 27, Last sequence update)
DE 05-JUL-2004 (TEMBLrel. 27, Last annotation update)
DB Adrenomedullin-2.
GN Name=ADM2;
OS Rugu rubripes (Japanese pufferfish) (Takifugu rubripes).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Actinopterygii; Neopterygii; Teleostei; Euteleostei; Neoteleostei;
OC Acanthomorphia; Acanthopterygii; Percomorpha; Tetraodontiformes;
OC Tetraodontidae; Tetraodontidae; Takifugu.
RN NCBI_TaxID=31033;
RP SEQUENCE FROM N.A.
RX MEDLINE=22984567; PubMed=14623291; DOI=10.1016/j.bbrcc.2003.10.111;
RA Ogoishi M., Inoue K., Takei Y.,
RT Identification of a novel adrenomedullin gene family in teleost
RT fish.";
RL Biochem. Biophys. Res. Commun. 311:1072-1077(2003).
DR EMBL: AB120296; BAD02342.1; -
SQ SEQUENCE 168 AA; 18544 MW; A5905E7CC112669B CRC64;

QY 1 TQQLRVCVLTGTCOVONLSHRLMQLMGAPGRDSPA PVPSPSPHSY 47
DB 121 SHQQLMKVACVLTGTCOVONLSHRLMQLMGAPGRDSPA PVPSPSPHSY 167

RESULT 6
Q61FS9 PRELIMINARY; PRT; 140 AA.
AC Q61FS9;
DT 05-JUL-2004 (TEMBLrel. 27, Created)
DT 05-JUL-2004 (TEMBLrel. 27, Last sequence update)
DE 05-JUL-2004 (TEMBLrel. 27, Last annotation update)
DB Intermedin (Fragment).
GN Name=IMDN;
OS Brachydanio rerio (Zebrafish) (Danio rerio).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Actinopterygii; Neopterygii; Teleostei; Ostariophysi; Cypriniformes;
OC Cyprinidae; Danio.
RN NCBI_TaxID=7955;
RP SEQUENCE FROM N.A.
RX PubMed=14615490; DOI=10.1074/jbc.M305332200;
RA Roh J., Chang C.L., Bhalla A., Klein C., Hsu S.Y.T.;
RT Intermedin is a calcitonin/CGRP family peptide acting through the

RT CRLR/RAMP receptor complexes.";
 RL J. Biol. Chem. 0:0-0(2003).
 RN [2]
 RP SEQUENCE FROM N.A.
 RA Hsu S.Y.T.;
 RL Submitted (NOV-2003) to the EMBL/GenBank/DBJ databases.
 CC -1- MISCELLANEOUS: The sequence shown here is derived from an
 DR EMBL/GenBank/DBJ third party annotation (TPA) entry.
 FT EMBL; BK004089; DAA0455.1; -
 SQ SEQUENCE 140 AA; 15816 MW; 509F74908CDD75D CRC64;
 Query Match 73.0%; Score 185.5; DB 2; Length 140;
 Best Local Similarity 77.3%; Pred. No. 7.2e-16;
 Matches 34; Conservative 7; Mismatches 2; Indels 1; Gaps 1;

QY 4 QQLRVCCVGTCCVQVONLSHRLMQLMGPAGRODSAPVDPSSPHSY 47
 Db, 97 QLMRVGCVIGTCOVONLSHRLMQLMGPAGRODSAPVDPSSPHSY 139

RESULT 7
 ID Q75XM8 PRELIMINARY; PRT; 174 AA.
 AC Q75XM8;
 DT 05-JUL-2004 (TREMBlrel. 27, Created)
 DT 05-JUL-2004 (TREMBlrel. 27, Last sequence update)
 DT 05-JUL-2004 (TREMBlrel. 27, Last annotation update)
 DE Adrenomedullin-1.
 GN Name=ADM1;
 OS Fugu rubripes (Japanese pufferfish) (Takifugu rubripes).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Actinopterygii; Neopterygii; Teleostei; Euteleostei; Neoteleostei;
 OC Acanthomorpha; Acanthopterygii; Percomorpha; Tetraodontiformes;
 OC Tetraodontidae; Tetraodontidae; Takifugu.
 RN NCB1_Taxid=31033;
 RP SEQUENCE FROM N.A.
 RX MEDLINE=22984567; PubMed=14623291; DOI=10.1016/j.bbrc.2003.10.111;
 RA Ogoehi M., Inoue K., Takei Y.;
 RT "Identification of a novel adrenomedullin gene family in teleost fish."
 RL Biochem. Biophys. Res. Commun. 311:1072-1077(2003).
 DR EMBL; AB120295; BAD02341.1; -
 DR GO; GO:0005576; C:extracellular; IEA.
 DR GO; GO:0005179; F:hormone activity; IEA.
 SQ SEQUENCE 174 AA; 20222 MW; 61535E41FCF8BD4D CRC64;

Query Match 34.4%; Score 87.5; DB 2; Length 174;
 Best Local Similarity 53.8%; Pred. No. 0.0038;
 Matches 21; Conservative 3; Mismatches 14; Indels 1; Gaps 1;

QY 9 GCVLGTCVQVONLSHRLMQLMGPAGRODSAPVDPSSPHSY 47
 Db 95 GCSLGTCTVHDLAFLRLHQL-GFOYKIDIAVPDKISPHGY 132

RESULT 8
 ID Q75XM4 PRELIMINARY; PRT; 123 AA.
 AC Q75XM4;
 DT 05-JUL-2004 (TREMBlrel. 27, Created)
 DT 05-JUL-2004 (TREMBlrel. 27, Last sequence update)
 DT 05-JUL-2004 (TREMBlrel. 27, Last annotation update)
 DE Adrenomedullin-5.
 GN Name=ADM5;
 OS Fugu rubripes (Japanese pufferfish) (Takifugu rubripes).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Actinopterygii; Neopterygii; Teleostei; Euteleostei; Neoteleostei;
 OC Acanthomorpha; Acanthopterygii; Percomorpha; Tetraodontiformes;
 OC Tetraodontidae; Tetraodontidae; Takifugu.
 RN NCB1_Taxid=31033;

RP SEQUENCE FROM N.A.
 RX MEDLINE=22984567; PubMed=14623291; DOI=10.1016/j.bbrc.2003.10.111;
 RA Ogoehi M., Inoue K., Takei Y.;
 RT "Identification of a novel adrenomedullin gene family in teleost fish."
 RL Biochem. Biophys. Res. Commun. 311:1072-1077(2003).
 DR EMBL; AB120299; BAD02345.1; -
 SQ SEQUENCE 123 AA; 13538 MW; D3ED4CDBCF4CEB CRC64;

Query Match 32.7%; Score 83; DB 2; Length 123;
 Best Local Similarity 41.0%; Pred. No. 0.0098;
 Matches 16; Conservative 9; Mismatches 12; Indels 2; Gaps 1;

QY 9 GCVLGTCVQVONLSHRLMQLMGPAGRODSAPVDPSSPHSY 47
 Db 85 GCQVGTCTVHDLANKLYQIGROGDESTRV--NDPQY 121

RESULT 9
 ID Q6L8K5 PRELIMINARY; PRT; 171 AA.
 AC Q6L8K5;
 DT 05-JUL-2004 (TREMBlrel. 27, Created)
 DT 05-JUL-2004 (TREMBlrel. 27, Last sequence update)
 DT 05-JUL-2004 (TREMBlrel. 27, Last annotation update)
 DE Preproadrenomedullin precursor.
 GN Name=prePROM;
 OS Cyprinus carpio (Common carp).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Actinopterygii; Neopterygii; Teleostei; Ostariophysi; Cypriniformes;
 OC Cyprinidae; Cyprinus.
 RN NCB1_Taxid=7962;
 RP SEQUENCE FROM N.A.
 RC TISSUE=Liver;
 RX PubMed=15242754; DOI=10.1016/j.ygscn.2004.05.002;
 RA Kono T., Sakai M.;
 RT "Molecular cloning and expression of preproadrenomedullin gene from common carp *Cyprinus carpio* L."
 RL Gen. Comp. Endocrinol. 138:78-88(2004).
 DR EMBL; AB120940; BAD19046.1; -
 DR GO; GO:0005576; C:extracellular; IEA.
 DR GO; GO:0005179; F:hormone activity; IEA.
 DR InterPro; IPR001710; Adrenomedullin.
 DR PRINTS; PR00801; ADRENOMEDULLIN.
 KW Signal.
 FT SIGNAL. 1 23 Potential.
 FT CHAIN 79 127 adrenomedullin.
 SQ SEQUENCE 171 AA; 19412 MW; A9595B9A11E5AC36 CRC64;

Query Match 30.9%; Score 78.5; DB 2; Length 171;
 Best Local Similarity 42.2%; Pred. No. 0.0053;
 Matches 19; Conservative 5; Mismatches 20; Indels 1; Gaps 1;

QY 3 AQLRVGCVLTCCVQVONLSHRLMQLMGPAGRODSAPVDPSSPHSY 47
 Db 84 SQSRSGCSLGTCTVHDLAFLRLHDLNNKL-KIGNAPAKKINFGY 127

RESULT 10
 ID ADML_HUMAN STANDARD; PRT; 185 AA.
 AC P35318;
 DT 01-FEB-1994 (Rel. 28, Created)
 DT 01-FEB-1994 (Rel. 28, Last sequence update)
 DT 05-JUL-2004 (Rel. 44, Last annotation update)
 DE ADM precursor [Contains: Adrenomedullin (AM); Proadrenomedullin N-20 terminal peptide (PROM-N20) (PROM N-terminal 20 peptide) (PAMP)].
 GN Name=ADM; Synonyms=AM;
 OS Homo sapiens (Human).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Primates; Catarrhini; Homidae; Homo.
 RN NCB1_Taxid=9606;

[1]
 RP SEQUENCE FROM N.A.
 RC TISSUE=Pheochromocytoma;
 RX MEDLINE=93343928; PubMed=7688224;
 RA Kitamura K., Sakata J., Kangawa K., Kojima M., Matsuo H., Eto T.;
 RT "Cloning and characterization of cDNA encoding a precursor for human
 RT adrenomedullin.";
 RL Biochem. Biophys. Res. Commun. 194:720-725(1993).
 RN [2]
 RP SEQUENCE FROM N.A.
 RC TISSUE=Liver;
 RX MEDLINE=94354869; PubMed=8074714;
 RA Ishihara T., Kojima M., Kangawa K., Hino J., Matsuo H.,
 RA Kitamura K., Eto T., Matsuo H.;
 RT "Genomic structure of human adrenomedullin gene.";
 RL Biochem. Biophys. Res. Commun. 203:631-639(1994).
 RN [3]
 RP SEQUENCE FROM N.A.
 RC TISSUE=Kidney;
 RX MEDLINE=22388257; PubMed=12477932; DOI=10.1073/pnas.242603899;
 RA Klausner R.D., Collins F.S., Wagner L., Shennan C.M., Schuler G.D.,
 RA Altschul S.F., Zeeberg B., Buetow K.H., Schaefer C.F., Bhat N.K.,
 RA Hopkins R.F., Jordan H., Moore T., Max S.I., Wang J., Hsieh P.,
 RA Diatchenko L., Marusina K., Farmer A.A., Rubin G.M., Hong L.,
 RA Stapleton M., Soares M.B., Bonaldo M.F., Casavant P.L., Scheetz T.E.,
 RA Brownstein M.J., Ustin T.B., Toehlyuk S., Carninci P., Prange C.;
 RA Raha S.S., Loggellano N.A., Peters G.J., Abramson R.D., Mulhany S.J.,
 RA Bosak S.A., McManus P.J., McKernan K.J., Malek J.A., Gunaratne P.H.,
 RA Villalón D.K., Muzny D.M., Sodergren E.J., Lu X., Gibbs R.A.,
 RA Fahy J., Helton E., Kettman M., Madan A., Rodriguez S., Sanchez A.,
 RA Whiting M., Madan A., Young A.C., Shevchenko Y., Bouffard G.G.,
 RA Blakesley R.W., Touchman J.W., Green E.D., Dickson M.C.,
 RA Rodriguez A.C., Grimwood J., Schmutz J., Myers R.M.,
 RA Butterfield Y.S.N., Krzywicki M.I., Skalek U., Smallus D.E.,
 RA Scherch A., Schein J.E., Jones S.J.M., Marra M.A.;
 RT "Generation and initial analysis of more than 15,000 full-length human
 RT and mouse cDNA sequences.";
 RL Proc. Natl. Acad. Sci. U.S.A. 99:16899-16903(2002).
 RN [4]
 RP SEQUENCE OF 95-146.
 RC TISSUE=Pheochromocytoma;
 RX MEDLINE=93249425; PubMed=8387282;
 RA Kitamura K., Kangawa K., Kawamoto M., Ichiki Y., Nakamura S.,
 RA Matsuo H., Eto T.;
 RT "Adrenomedullin: a novel hypotensive peptide isolated from human
 RT pheochromocytoma.";
 RL Biochem. Biophys. Res. Commun. 192:553-560(1993).
 RN [5]
 RP REVIEW.
 RX MEDLINE=98240137; PubMed=9578982; DOI=10.1006/erne.1998.0164;
 RA Samson W.K.;
 RT "Proadrenomedullin-derived peptides.";
 RL Front. Neuroendocrinol. 19:100-127(1998).
 RN [6]
 RP REVIEW.
 RX MEDLINE=20053666; PubMed=10588445; DOI=10.1016/S0167-0115(99)00025-7;
 RA Champion H.C., Nussdorfer G.G., Kadowitz P.J.;
 RT "Structure-activity relationships of adrenomedullin in the circulation
 RT and adrenal gland.";
 RL Regul. Pept. 85:1-8(1999).
 CC -1- FUNCTION: AM and PAMP are potent hypotensive and vasodilator
 CC agents. Numerous actions have been reported most related to the
 CC physiologic control of fluid and electrolyte homeostasis. In the
 CC kidney, am is diuretic and natriuretic, and both am and pamp
 CC inhibit aldosterone secretion by direct adrenal actions. In
 CC pituitary gland, both peptides at physiologically relevant doses
 CC inhibit basal ACTH secretion. Both peptides appear to act in brain
 CC and pituitary gland to facilitate the loss of plasma volume,
 CC actions which complement their hypotensive effects in blood
 CC vessels.
 CC -1- SUBCELLULAR LOCATION: Secreted.

CC -1- TISSUE SPECIFICITY: Highest levels found in pheochromocytoma and
 CC adrenal medulla. Also found in lung, ventricle and kidney tissues.
 CC -1- SIMILARITY: Belongs to the adrenomedullin family.
 CC -----
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 CC -----
 CC EMBL; D14874; BAA03589.1; -
 CC EMBL; S73906; AAC60642.1; -
 CC EMBL; BC015861; AAH15861.1; -
 CC EMBL; D43639; BAA07756.1; ALT_SEQ.
 CC PIR; JC2351; JN0684.
 CC Genem; HGNC:259; ADM.
 CC H-InvDB; HIK0009441; -
 CC MIM; 103275; -
 CC DR GO; GO:0005615; Cytoplasmic space; TAS.
 CC DR GO; GO:0005625; Cytoplasmic fraction; TAS.
 CC DR GO; GO:0005102; F-actin binding; TAS.
 CC DR GO; GO:0006171; P-cAMP biosynthesis; TAS.
 CC DR GO; GO:0007267; P-cAMP biosynthesis; TAS.
 CC DR GO; GO:0008015; P-circulation; TAS.
 CC DR GO; GO:0007565; P-pregnancy; TAS.
 CC DR GO; GO:0006701; P-progesterone biosynthesis; TAS.
 CC DR GO; GO:0009611; P-response to wounding; TAS.
 CC DR GO; GO:0007165; P-signal transduction; TAS.
 CC DR InterPro; IPR001710; Adrenomedullin.
 CC DR Pfam; PF02039; Adrenomedullin; 1.
 CC DR PRINTS; PRO0801; ADRENOMEDULLIN.
 CC KW Adrenomedullin; Cleavage on pair of basic residues;
 CC Direct protein sequencing; Hormone; Polymorphism; signal.
 CC FT SIGNATURE 2 21
 CC FT PEPTIDE 22 41
 CC FT PROPEP 45 92
 CC FT PROPEP 148 185
 CC FT DISULFID 110 115
 CC FT MOD_RES 41 41
 CC FT MOD_RES 146 146
 CC FT VARIANT 50 50
 CC FT VARIANT 50 50
 CC FT SEQUENCE 185 AA; 20420 MW; 64C7D2A0B4654DFB CRC64;
 CC Query Match 28.5%; Score 72.5; DB 1; Length 185;
 CC Best Local Similarity 38.5%; Pred. No. 0.34; Indels 1; Gaps 1;
 CC Matches 15; Conservative 6; Mismatches 17;
 CC Db 109 GCVLGTGVONTSLRMLMGAPGRODSAPVDPSPSPHY 47
 CC 9 GCVLGTGVONTSLRMLMGAPGRODSAPVDPSPSPHY 47
 CC 109 GCVLGTGVONTSLRMLMGAPGRODSAPVDPSPSPHY 47
 CC RESULT 11
 CC ADML_MOUSE STANDARD; PRT; 184 AA.
 CC AC P97257; P97453; -
 CC DT 16-OCT-2001 (Rel. 40, Created)
 CC DT 16-OCT-2001 (Rel. 40, Last sequence update)
 CC DT 05-JUL-2004 (Rel. 44, Last annotation update)
 CC DE ADM precursor [Contains: Adrenomedullin (ADM); Proadrenomedullin N-20
 CC terminal peptide (ProADM-N20) (ProADM N-terminal 20 peptide) (PAMP)].
 CC GN Name-Adm.
 CC OS Mus musculus (Mouse).
 CC OC Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;
 CC OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
 CC OX NCBI_TaxID=10090;
 CC RN [1]

RP SEQUENCE FROM N.A.
 RC STRAIN=129/SV;
 RX MEDLINE=97092892; PubMed=8938454; DOI=10.1006/geno.1996.0576;
 RA Okazaki T., Ogawa Y., Tamura N., Mori Y., Isse N., Aoki T.,
 RA Rochelle J.M., Takeo M.M., Seidin M.F., Nakao K.,
 RT "Genomic organization, expression, and chromosomal mapping of the
 RT mouse adrenomedullin gene.";
 RL Genomics 37:395-399(1996).
 [2]
 RP SEQUENCE FROM N.A.
 RC STRAIN=C57BL/6J;
 RX MEDLINE=9904675; PubMed=9808778; DOI=10.1006/dbio.1998.9073;
 RA Yotsumoto S., Shimada T., Chai C.Y., Nakashima H., Fujiwara H.,
 RA Ko M.S.H.;
 RT "Expression of adrenomedullin, a hypotensive peptide, in the
 RT trophoblast giant cells at the embryo implantation site in mouse.";
 RL Dev. Biol. 203:264-275(1998).
 CC -1- FUNCTION: AM and PAMP are potent hypotensive and vasodilator
 CC agents.
 CC -1- SUBCELLULAR LOCATION: Secreted.
 CC -1- SIMILARITY: Belongs to the adrenomedullin family.
 CC -----
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 CC -----
 CC EMBL; D78349; BAA11367.1; -;
 DR EMBL; U77630; AAB36535.1; -;
 DR MGD; MGI:108058; Adm.
 DR InterPro; IPR001710; Adrenomedullin.
 DR Pfam; PF02039; Adrenomedullin; 1.
 DR PRINTS; PRO0801; ADRENOMEDULLIN.
 KW Amidation; Cleavage on pair of basic residues; Hormone; Signal.
 FT SIGNAL 1 21
 FT PROPEP 22 41 By similarity.
 FT PROPEP 45 92 Proadrenomedullin N-20 terminal peptide.
 FT PROPEP 95 144 By similarity.
 FT PROPEP 121 184 Adrenomedullin.
 FT PROPEP 121 184 PreproAM C-terminal fragment (By
 FT similarity).
 FT DISULFID 108 113 Arginine amide (G-42 provides amide
 FT MOD_RES 41 41 By similarity.
 FT MOD_RES 144 144 Tyrosine amide (G-145 provides amide
 FT MOD_RES 144 144 group) (By similarity).
 FT CONFLICT 173 173 A -> G (in Ref. 2).
 FT SEQUENCE 184 AA; 20764 MW; C88C99045A79C898 CRC64;
 SQ
 Query Match 28.1%; Score 71.5; DB 1; Length 184;
 Best Local Similarity 38.5%; Pred. No. 0.46; Mismatches 1; Gaps 1;
 Matches 15; Conservative 6; Indels 1;
 Oy 9 GCVLGTGVONLSHRLMQLMGPAGRODSAPVDPSPSHSY 47
 Db 107 GCRFCTGVOKLAHQIYQLT-DKDDGMAPRYKISPPQGI 144

CC Bovinae; Bos.
 OX NCBI_TaxID=9913;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC TISSUE=Aorta;
 RX MEDLINE=98244567; PubMed=9585168; DOI=10.1016/S0024-3205(98)00079-4;
 RA Barker S., Wood E., Clark A.J.L., Corder R.,
 RT "Cloning of bovine preproadrenomedullin and inhibition of its basal
 RT expression in vascular endothelial cells by staurosporine.";
 RL Life Sci. 62:1407-1415(1998).
 CC -1- FUNCTION: Hypotensive peptide. May function as a hormone in
 CC circulation control (By similarity).
 CC -1- SUBCELLULAR LOCATION: Secreted.
 CC -1- SIMILARITY: Belongs to the adrenomedullin family.
 CC -----
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 CC or send an email to license@isb-sib.ch).
 CC -----
 CC EMBL; AJ001613; CA04866.1; -;
 DR InterPro; IPR001710; Adrenomedullin.
 DR Pfam; PF02039; Adrenomedullin; 1.
 DR PRINTS; PRO0801; ADRENOMEDULLIN.
 KW Amidation; Cleavage on pair of basic residues; Hormone; Signal.
 FT SIGNAL 1 21
 FT PROPEP 22 41 By similarity.
 FT PROPEP 45 92 Proadrenomedullin N-20 terminal peptide.
 FT PROPEP 95 146 By similarity.
 FT PROPEP 148 188 Adrenomedullin.
 FT PROPEP 148 188 PreproAM C-terminal fragment (By
 FT similarity).
 FT DISULFID 110 115 Arginine amide (G-42 provides amide
 FT MOD_RES 41 41 By similarity).
 FT MOD_RES 146 146 Tyrosine amide (G-147 provides amide
 FT MOD_RES 146 146 group) (By similarity).
 FT SEQUENCE 188 AA; 20981 MW; 3002F79AB3B612C CRC64;
 SQ
 Query Match 28.1%; Score 71.5; DB 1; Length 188;
 Best Local Similarity 38.5%; Pred. No. 0.47; Mismatches 18; Indels 1;
 Matches 15; Conservative 5; Gaps 1;
 Oy 9 GCVLGTGVONLSHRLMQLMGPAGRODSAPVDPSPSHSY 47
 Db 109 GCRFCTGVOKLAHQIYHFT-DKDDGSAFPRKISPPQGI 146

RESULT 12
 ADML BOVIN STANDARD; PRT; 188 AA.
 AC 062827;
 DT 16-OCT-2001 (Rel. 40, Created)
 DT 16-OCT-2001 (Rel. 40, Last sequence update)
 DT 05-JUL-2004 (Rel. 44, Last annotation update)
 DE ADM precursor [Contains: Adrenomedullin (AM); Proadrenomedullin N-20
 DE terminal peptide (ProAM-N20) (ProAM N-terminal 20 peptide) (PAMP)].
 GN Name=ADM;
 OS Bos taurus (Bovine).
 OC Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Cetartiodactyla; Ruminantia; Pecora; Bovidae;

RESULT 13
 ID 095KP0 PRELIMINARY; PRT; 188 AA.
 AC 095KP0;
 DT 01-DEC-2001 (TREMBlrel. 19, Created)
 DT 01-DEC-2001 (TREMBlrel. 19, Last sequence update)
 DT 01-MAR-2004 (TREMBlrel. 26, Last annotation update)
 DE Adrenomedullin.
 GN Name=PBAM-2;
 OS Bos taurus (Bovine).
 OC Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Cetartiodactyla; Ruminantia; Pecora; Bovidae;
 OC Bovinae; Bos.
 OX NCBI_TaxID=9913;
 RN [1]
 RP SEQUENCE FROM N.A.
 RX MEDLINE=21630318; PubMed=11754956; DOI=10.1016/S0196-9781(01)00529-0;
 RA Kitamura K., Matsui E., Kato J., Katoh F., Kita T., Tsuji T.,
 RA Kangawa K., Eto T.;
 RT "Adrenomedullin (11-26): a novel endogenous hypertensive peptide
 RT isolated from bovine adrenal medulla.";
 RL Peptides 22:1713-1718(2001).
 DR EMBL; AB055107; BAB62176.1; -;

DR GO:0005576; C:extracellular; IEA.
 DR GO:0005179; F:hormone activity; IEA.
 DR InterPro: IPR001710; Adrenomedullin.
 DR Pfam: PF02039; Adrenomedullin; 1.
 DR PRINTS: PR00801; ADRENOMEDULLIN.
 SQ SEQUENCE 188 AA; 20963 MW; 6102869A756DCAB6 CRC64;

Query Match 28.1%; Score 71.5; DB 2; Length 188;
 Best Local Similarity 38.5%; Pred. No. 0.47;
 Matches 15; Conservative 5; Mismatches 18; Indels 1; Gaps 1;

QY 9 GCVLGTCQVONLSHRLWQLMGPAGRODSAPVDPSSPSHY 47
 Db 109 GCRFGTCVQKLAHQIYQFT-DKDKGASAPRSKISPGY 146

RESULT 14
 ID ADML_PIG STANDARD; PRT; 188 AA.

AC P53366;
 DT 01-OCT-1996 (Rel. 34, Last sequence update)
 DT 01-OCT-1996 (Rel. 34, Last sequence update)
 DT 05-JUL-2004 (Rel. 44, Last annotation update)
 DE ADM precursor [Contains: Adrenomedullin (AM); Proadrenomedullin N-20 terminal peptide (ProAM-N20) (ProAM N-terminal 20 peptide) (PAMP)].
 GN Name=ADM; Synonyms=AM;
 OS Sus scrofa (Pig).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Cetartiodactyla; Suina; Suidae; Sus.
 NCBI_TaxID=9823;
 RX MEDLINE=94139945; PubMed=8043068; DOI=10.1016/0014-5793(94)80289-0;
 RA Kitamura K., Kangawa K., Kojima M., Ichiki Y., Matsuo H., Eto T.;
 RT "Complete amino acid sequence of porcine adrenomedullin and cloning of
 RT cDNA encoding its precursor."
 RL FEBS Lett. 338:306-310(1994).

RP SEQUENCE FROM N.A.
 RC TISSUE=Adrenal medulla;
 RA MEDLINE=94139945; PubMed=8043068; DOI=10.1016/0014-5793(94)80289-0;
 RA Kitamura K., Kangawa K., Kojima M., Ichiki Y., Matsuo H., Eto T.;
 RT "Complete amino acid sequence of porcine adrenomedullin and cloning of
 RT cDNA encoding its precursor."
 RL FEBS Lett. 338:306-310(1994).

RT SEQUENCE OF 22-41.
 RC TISSUE=Adrenal medulla;
 RA MEDLINE=94139945; PubMed=8043068; DOI=10.1016/0014-5793(94)80289-0;
 RA Kitamura K., Kangawa K., Kojima M., Ichiki Y., Matsuo H., Eto T.;
 RT "Complete amino acid sequence of porcine adrenomedullin and cloning of
 RT cDNA encoding its precursor."
 RL FEBS Lett. 338:306-310(1994).

CC -1- SUBCELLULAR LOCATION: Secreted.
 CC -1- TISSUE SPECIFICITY: Highly expressed in adrenal glands, lung and
 CC kidney.
 CC -1- SIMILARITY: Belongs to the adrenomedullin family.

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DR EMBL: D14875; BA03590.1; -.
 DR PIR: S41600; S41600.
 DR InterPro: IPR001710; Adrenomedullin.
 DR InterPro: IPR01038; Calycin.
 DR Pfam: PF02039; Adrenomedullin; 1.
 DR PRINTS: PR00801; ADRENOMEDULLIN.
 KM Amidation; Cleavage on pair of basic residues;
 KM Direct protein sequencing; Hormone; Signal.
 FT SIGNAL 1 21
 FT PEPIDE 22 92 Proadrenomedullin N-20 terminal peptide.
 FT PROPEP 45 92

FT PEPTIDE 95 146 Adrenomedullin.
 FT PROPEP 153 188 PreproAM C-terminal fragment (By
 FT DLSUFID 110 115 similarity).
 FT MOD_RES 41 41 Arginine amide (G-42 provides amide
 FT MOD_RES 146 146 Tyrosine amide (G-147 provides amide
 FT MOD_RES 146 146 group).
 SQ SEQUENCE 188 AA; 20893 MW; 71749460F5660A61 CRC64;

Query Match 27.8%; Score 70.5; DB 1; Length 188;
 Best Local Similarity 38.5%; Pred. No. 0.63;
 Matches 15; Conservative 5; Mismatches 18; Indels 1; Gaps 1;

QY 9 GCVLGTCQVONLSHRLWQLMGPAGRODSAPVDPSSPSHY 47
 Db 109 GCRFGTCVQKLAHQIYQFT-DKDKGASAPRSKISPGY 146

RESULT 15
 ID Q7S575 PRELIMINARY; PRT; 927 AA.

AC Q7S575;
 DT 01-MAR-2004 (TrEMBLrel. 26, Created)
 DT 01-MAR-2004 (TrEMBLrel. 26, Last sequence update)
 DT 01-MAR-2004 (TrEMBLrel. 26, Last annotation update)
 DE Hypothetical protein.
 GN Name=NCU02295.1;
 OS Neurospora crassa.
 OC Eukaryota; Fungi; Ascomycota; Pezizomycotina; Sordariomycetes;
 OC Sordariomycetidae; Sordariales; Sordariaceae; Neurospora.
 NCBI_TaxID=5141;
 RN [1]

RP SEQUENCE FROM N.A.
 RC STRAIN=OR74A;
 RA Galagan J.E., Calvo S.E., Borkovich K.A., Selker E.U., Read N.D.,
 RA Jaffe D., Fitzhugh W., Ma L.-J., Smirnov S., Purcell S., Rehm B.,
 RA Elkins T., Engels R., Wang S., Nielsen C.B., Butler J., Endrizzi M.,
 RA Qui D., Ianakiev P., Pedersen D., Nelson M., Washburne M.,
 RA Selitrenikoff C.P., Kinsey J.A., Braun E.L., Zelter A., Schulte U.,
 RA Kothe G.O., Jedd G., Mewes W., Staben C., Marcotte E., Greenberg D.,
 RA Roy A., Foley K., Naylor J., Thoman N., Barrett R., Gierke S.,
 RA Kamal M., Kamysseis M., Mauceli E., Bielke C., Rudd S., Prihman D.,
 RA Krystofova S., Raamsen C., Metzberg R.L., Perkins D.D., Kroken S.,
 RA Cogoni C., Macino G., Cacheseide D., Li W., Pratt R.J., Omani S.A.,
 RA Desouza C.C., Glaes L., Orbach M.J., Berglund J., Volker R.,
 RA Yarden O., Plamann M., Seiler S., Dunlap J., Radford A., Aramayo R.,
 RA Natvig D.O., Alex L.A., Mannhaupt G., Ebbold D.J., Freitag M.,
 RA Paulsen I., Sachs M.S., Lander E.S., Nusbaum C., Birren B.;
 RT "The Genome Sequence of the Filamentous Fungus Neurospora crassa."
 RL Nature 010-0(2003).

CC -1- CAUTION: The sequence shown here is derived from an
 CC EMBL/GenBank/DBD whole genome shotgun (WGS) entry which is
 CC preliminary data.

DR EMBL: AABX01000355; EAA30703.1; -.
 DR HSSP: P78356; 1B01.
 DR GO:0016308; F:1-phosphatidylinositol-4-phosphate 5-kinase. . .; IEA.
 DR InterPro: IPR002498; PI3K.
 DR Pfam: PF01504; PI3PK; 1.
 KM Hypothetical protein.
 SQ SEQUENCE 927 AA; 102558 MW; 21PDC49FA419E932 CRC64;

Query Match 27.6%; Score 70; DB 2; Length 927;
 Best Local Similarity 72.2%; Pred. No. 4.2;
 Matches 13; Conservative 1; Mismatches 4; Indels 0; Gaps 0;

QY 30 PACGDSAPVDPSSPSHY 47
 Db 123 PACGRDSAPVDPSSPSHY 140

Search completed: May 4, 2005, 18:14:12
 Job time : 178 secs

GenCore version 5.1.6
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OM protein - protein search, using sw model

Run on: May 4, 2005, 17:58:05 ; Search time 39 Seconds
(Without alignments)
115.954 Million cell updates/sec

Title: US-10-723-368-5

Sequence: 1 TQQLRVCCVLCGTQCVQNL.....MGPGRQDSAPVDPSSPHSY 47

Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 283416 seqs, 96216763 residues

Total number of hits satisfying chosen parameters: 283416

Minimum DB seq length: 0
Maximum DB seq length: 200000000

Post-processing: Minimum Match 0%
Maximum Match 100%

Liecing first 45 summaries

Database :

1: PIR 79:.*
2: p1r1:.*
3: p1r2:.*
4: p1r3:.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length DB	ID	Description
1	72.5	28.5	185	2 JN0684	adrenomedullin pre
2	70.5	27.8	188	2 S41600	adrenomedullin - p
3	69.5	27.4	185	2 JN0766	adrenomedullin pre
4	58	22.8	339	2 AC3406	1-lactate permease
5	58	22.8	560	2 AC8731	alpha-1A adenylyl
6	58	22.8	1544	2 T29482	hypothetical prote
7	57	22.4	2	T02844	cdclc-related prot
8	54	21.3	283	2 E82662	ABC transporter AT
9	54	21.3	792	2 T26050	hypothetical prote
10	53.5	21.1	303	2 A43708	gamma-interferon-1
11	53.5	21.1	345	2 A37845	carboxylesterase (
12	53	20.9	136	2 T35335	hypothetical prote
13	53	20.9	192	2 AE2414	hypothetical prote
14	53	20.9	272	2 H95954	probable Sit2-like
15	53	20.9	351	2 E97187	cytochrome P450 4A6
16	53	20.9	450	2 S07051	cysteine proteinase
17	53	20.9	450	2 S12099	hypothetical prote
18	53	20.9	765	2 S74598	hypothetical prote
19	53	20.9	860	2 JC5702	ERBB kinase activa
20	53	20.9	868	2 JC5701	ERBB kinase activa
21	52.5	20.7	347	2 T26349	hypothetical prote
22	52.5	20.7	910	2 AE3380	valine-CRNA ligase
23	52	20.5	381	2 S38663	hypothetical prote
24	52	20.5	665	2 C81439	probable integral
25	52	20.5	2870	2 H96974	cyclic beta 1-2 gl
26	52	20.5	3898	1 GNMVHB	genome polyprotein
27	52	20.5	3898	1 S57437	genome polyprotein
28	52	20.5	3898	2 S58295	polyprotein - hog
29	51.5	20.3	461	2 B95388	Probable (EC 1.1.1

30	51.5	20.3	977	2 I52657	seizure-related pr
31	51	20.1	106	2 A05148	hypothetical reduc
32	51	20.1	407	2 I52703	42K membrane glyco
33	51	20.1	552	2 S15555	NAD synthase (EC 6
34	51	20.1	771	2 G71409	probable replicati
35	51	20.1	813	2 E85135	hypothetical prote
36	51	20.1	1359	2 T10235	xanthine dehydroge
37	51	20.1	3898	1 GNMVHC	genome polyprotein
38	51	20.1	4544	1 S02392	alpha-2-macroglobu
39	50.5	19.9	296	2 T12469	hypothetical prote
40	50.5	19.9	317	2 S22087	peroxidase (EC 1.1
41	50.5	19.9	397	2 T19022	hypothetical prote
42	50.5	19.9	460	2 AG2262	hypothetical prote
43	50.5	19.9	488	2 T31622	hypothetical prote
44	50.5	19.9	503	2 A85900	hypothetical prote
45	50.5	19.9	503	2 E91055	hypothetical prote

ALIGNMENTS

RESULT 1
JN0684
adrenomedullin precursor - human
C:Date: 03-Feb-1994 #sequence revision 03-Feb-1994 #text_change 09-Jul-2004
C:Accession: JC2351; JN0684; E0548; JN0476
R:Rihimatsu, T.; Kojima, M.; Kangawa, K.; Hino, J.; Matsuo, H.; Kitamura, K.; Eto, T.;
Biochem. Biophys. Res. Commun. 203, 631-639, 1994
A:Title: Genomic structure of human adrenomedullin gene.
A:Reference number: JC2351; PMID:94354869; PMID:8074714
A:Accession: JC2351
A:Molecule type: DNA
A:Residues: 1-185 <185>
A:Cross-references: UNIPROT:P35318; GB:S73906; NID:G765329; PIDN:AA06042.1; PID:G765330
A:Experimental source: pheochromocytoma
R:Kitamura, K.; Sakata, J.; Kangawa, K.; Kojima, M.; Matsuo, H.; Eto, T.
Biochem. Biophys. Res. Commun. 194, 720-725, 1993
A:Title: Cloning and characterization of cDNA encoding a precursor for human adrenomedullin.
A:Reference number: JN0684; PMID:93343928; PMID:7688224
A:Accession: JN0684
A:Molecule type: mRNA
A:Residues: 1-185 <185>
A:Cross-references: GB:D14874; NID:G455470; PIDN:BA03589.1; PID:G500612
A:Accession: E0548
A:Molecule type: protein
A:Residues: 22-41 <R12>
R:Kitamura, K.; Kangawa, K.; Kawamoto, M.; Ichiki, Y.; Nakamura, S.; Matsuo, H.; Eto, T.
Biochem. Biophys. Res. Commun. 192, 553-560, 1993
A:Title: Adrenomedullin: A novel hypotensive peptide isolated from human pheochromocytoma
A:Reference number: JN0476; PMID:93249425; PMID:8387282
A:Accession: JN0476
A:Molecule type: protein
A:Residues: 95-146 <R13>
A:Experimental source: pheochromocytoma
C:Genetics:
A:Gene: GDB:ADM
A:Cross-references: GDB:217070; OMIM:103275
A:Map position: 11pter-11qter
A:Insertions: 33/2; 83/2
C:Keywords: amidated carboxyl end; blood pressure control; hormone
F:1-21/Domain: signal sequence #status predicted <SIG>
F:22-185/Product: proadrenomedullin #status predicted <PEP>
F:95-146/Domain: proadrenomedullin amino-terminal 20 peptide #status predicted <PAP>
F:147-185/Domain: adrenomedullin #status experimental <MAT>
F:147-185/Domain: carboxyl-terminal propeptide #status predicted <CTP>
F:147-185/Domain: amidated carboxyl end (Arg) (amide in mature form from following glyc
F:110-115/Disulfide bonds: #status experimental
F:146/Modified site: amidated carboxyl end (Tyr) (amide in mature form from following gly)

Query Match 28.5%; Score 72.5; DB 2; Length 185;
Best Local Similarity 38.5%; Pred. No. 0.037;
Matches 15; Conservative 6; Mismatches 17; Indels 1; Gaps 1;

QY 9 GCVLGTCQVONTLSHRLWQMGAPAGRODSAPVDPSSPHSY 47
DB 109 GCRFGCTCTVQKLAHQIYQFT-DKDXDGNVAPRKSISPOGY 146

RESULT 2
S41600
adrenomedullin - pig
C/Species: Sus scrofa domestica (domestic pig)
C/Date: 19-Mar-1997 #sequence_revision 19-Mar-1997 #text_change 09-Jul-2004
C/Accession: S41600
R/Kitamura, K.; Kangawa, K.; Kojima, M.; Ichiki, Y.; Matsuo, H.; Eto, T.
FEBS Lett. 338, 306-310, 1994
A/Title: Complete amino acid sequence of porcine adrenomedullin and cloning of cDNA encod
A/Accession: S41600; MUID:94139945; PMID:8043068
A/Status: preliminary
A/Molecule type: mRNA
A/Residues: 1-188 <KIT>
A/Cross-references: UNIPROT:P53366; GB:D14875; NID:g439721; PIDN:BA03590.1; PID:g496379

Query Match 27.8%; Score 70.5; DB 2; Length 188;
Best Local Similarity 38.5%; Pred. No. 0.069;
Matches 15; Conservative 5; Mismatches 18; Indels 1; Gaps 1;

QY 9 GCVLGTCQVONTLSHRLWQMGAPAGRODSAPVDPSSPHSY 47
DB 109 GCRFGCTCTVQKLAHQIYQFT-DKDXDGNVAPRKSISPOGY 146

RESULT 3
JN0766
adrenomedullin precursor - rat
C/Species: Rattus norvegicus (Norway rat)
C/Date: 30-Sep-1993 #sequence_revision 20-Aug-1994 #text_change 09-Jul-2004
C/Accession: JN0766; PNO610
R/Sakakura, J.; Shimokubo, T.; Kitamura, K.; Nakamura, S.; Kangawa, K.; Matsuo, H.; Eto, T.
Biochem. Biophys. Res. Commun. 195, 921-927, 1993
A/Title: Molecular cloning and biological activities of rat adrenomedullin, a hypotensive
A/Reference number: JN0766; MUID:93384621; PMID:7690563
A/Accession: JN0766
A/Molecule type: mRNA
A/Residues: 1-185 <SAK>
A/Cross-references: UNIPROT:P43145
A/Accession: PNO610
A/Molecule type: protein
A/Residues: 22-41 <SA2>
C/Comment: This precursor contains a unique 20-amino acid sequence designated proadrenom
essure control.
C/Keywords: amidated carboxyl end
F/1-21/Domain: signal sequence #status predicted <SIG>
F/22-185/Product: proadrenomedullin #status predicted <PEU>
F/94-143/Product: adrenomedullin amino-terminal 20 peptide #status predicted <PAF>
F/41/Modified site: amidated carboxyl end (Arg) (amide in mature form from following g1
F/143/Modified site: amidated carboxyl end (Tyr) (amide in mature form from following g1

Query Match 27.4%; Score 69.5; DB 2; Length 185;
Best Local Similarity 35.9%; Pred. No. 0.092;
Matches 14; Conservative 7; Mismatches 17; Indels 1; Gaps 1;

QY 9 GCVLGTCQVONTLSHRLWQMGAPAGRODSAPVDPSSPHSY 47
DB 106 GCRFGCTCTVQKLAHQIYQFT-DKDXDGNVAPRKSISPOGY 143

RESULT 4
AC3406
1-lactate permease [imported] - Brucella melitensis (strain 16M)
C/Species: Brucella melitensis
C/Date: 01-Feb-2002 #sequence_revision 01-Feb-2002 #text_change 09-Jul-2004
C/Accession: AC3406

R,DeIvecchio, V.G.; Kapral, V.; Redkar, R.J.; Patra, G.; Mujar, C.; Los, T.; Ivanova, P.
; Mazur, M.; Goldsman, E.; Selkov, E.; Elizer, P.H.; Haglue, S.; O'Callaghan, D.; Letesec
Proc. Natl. Acad. Sci. U.S.A. 99, 443-448, 2002
A/Title: The genome sequence of the facultative intracellular pathogen Brucella melitensis
A/Reference number: AD3252; PMID:11756688
A/Accession: AC3406
A/Status: preliminary
A/Molecule type: DNA
A/Residues: 1-339 <KUR>
A/Cross-references: UNIPROT:O8YGC6; GB:AE008917; PIDN:AAL52414.1; PID:g17983216; GSPDB:GR
A/Experimental source: strain 16M
C/Genetics:
A/Gene: BMEI233
A/Map position: I

Query Match 22.8%; Score 58; DB 2; Length 339;
Best Local Similarity 27.7%; Pred. No. 5.9;
Matches 13; Conservative 6; Mismatches 20; Indels 8; Gaps 1;

QY 8 VECVLGTCQVONTLSHRLWQMGAPAGRODSAPVDPSSPHS 46
DB 31 VDVIATCSMALAFRLFQPRRTWTSTGKEBTNAPVQPRSHS 77

RESULT 5
A38731
alpha-1A adrenergic receptor - rat
C/Species: Rattus norvegicus (Norway rat)
C/Date: 22-Jan-1993 #sequence_revision 22-Jan-1993 #text_change 09-Jul-2004
C/Accession: A38731; A53280
R/Lomansney, J.W.; Cotecchia, S.; Lorenz, W.; Leung, W.Y.; Schwinn, D.A.; Yang-Feng, T.L.;
J. Biol. Chem. 266, 6365-6369, 1991
A/Title: Molecular cloning and expression of the cDNA for the alpha-1A-adrenergic recept
A/Reference number: A38731; MUID:91177889; PMID:1706716
A/Accession: A38731
A/Molecule type: mRNA
A/Residues: 1-560 <LOM>
A/Cross-references: UNIPROT:P23944; GB:M60654; NID:g202761; PIDN:AAA63477.1; PID:g202762
R/Perez, D.M.; Pisacit, M.T.; Graham, R.M.
Mol. Pharmacol. 40, 876-883, 1991
A/Title: Solution-phase library screening for the identification of rare clones: isolatic
A/Reference number: A53280; MUID:92100054; PMID:1661838
A/Accession: A53280
A/Status: preliminary; not compared with conceptual translation
A/Molecule type: mRNA
A/Residues: 1-36, 'P', 38-58, 'I', 60-202, 'I', 204-305, 'R', 307-366, 'I', 368-370, 'I', 372-559 <PE
A/Experimental source: hippocampus
A/Note: sequence extracted from NCBI backbone (NCBIP:73541)
C/Superfamily: vertebrate rhodopsin
C/Keywords: G protein-coupled receptor; glycoprotein; transmembrane protein

Query Match 22.8%; Score 58; DB 2; Length 560;
Best Local Similarity 30.9%; Pred. No. 10;
Matches 17; Conservative 3; Mismatches 15; Indels 20; Gaps 2;

QY 4 QLRVGCVLGTCQVONTLSHRLWQMGAPAGROD-----SAPVDPSSPH 45
DB 409 RLNR-----CQRRRRRLRLMSLRPLASLDNRARRLRPPSHSPKPPSSPH 456

RESULT 6
T29482
hypothetical protein F08B4.2 - Caenorhabditis elegans
C/Species: Caenorhabditis elegans
C/Date: 15-Oct-1999 #sequence_revision 15-Oct-1999 #text_change 09-Jul-2004
C/Accession: T29482
R/Stelljes, L.; Gattung, S.
submitted to the EMBL Data Library, March 1996
A/Description: The sequence of C. elegans cosmid F08B4.
A/Reference number: Z20625
A/Accession: T29482
A/Status: preliminary; translated from GB/EMBL/DDBJ
A/Molecule type: DNA

A;Residues: 1-1544 <STES>
A;Cross-references: UNIPROT:Q019194; EMBL:U52002; PDB:1AAB37728.1; GSPDB:GN00022; CESP:FC
A;Experimental source: strain Bristol N2; clone F08B4
C;Genetics:
A;Gene: CESP:F08B4.2
A;Map position: 4
A;Intons: 25/3; 74/3; 147/3; 238/1; 290/2; 391/2; 452/3; 526/3; 670/3; 734/1; 779/3; 83

Query Match 22.4%; Score 58; DB 2; Length 1544;
Best Local Similarity 25.4%; Pred. No. 30;
Matches 15; Conservative 7; Mismatches 23; Indels 14; Gaps 1;

Qy 3 AQLRVGCVLTGTCOV-----ONTLSRLMQLMGPRQDSAPVDPSSPSHY 47
Db 1275 AVTVAGATIGICAVCFMGRYKTAQRNANSHSYQKGPYVHPMTMGVDPRTEYDY 1333

RESULT 7
T02844
cdc16-related protein, L3169.1 [imported] - Leishmania major (strain Friedlin)
C;Date: 24-Mar-1999 #sequence_revision 24-Mar-1999 #text_change 09-Jul-2004
C;Accession: H81461; T02844
R;Wylet, P.J.; Audleman, L.; devos, T.; Hixson, G.; Kiser, P.; Lemley, C.; Magness, C.;
Proc. Natl. Acad. Sci. U.S.A. 96, 2902-2906, 1999
A;Title: Leishmania major Friedlin chromosome 1 has an unusual distribution of protein-c
A;Reference number: A81455; MUID:99178987; PMID:10077609
A;Accession: H81461
A;Status: preliminary
A;Molecule type: DNA
A;Residues: 1-1784 <PVL>
A;Cross-references: UNIPROT:Q94606; GB:AE001274; NID:G3264850; PDB:1AAC24667.1; PID:G326
A;Experimental source: strain MHOM/IL/81/Friedlin
C;Genetics:
A;Gene: L3169.1
A;Map position: 1

Query Match 22.4%; Score 57; DB 2; Length 1784;
Best Local Similarity 52.6%; Pred. No. 47;
Matches 10; Conservative 5; Mismatches 4; Indels 0; Gaps 0;

Qy 16 QVONLSHRLMQLMGPRQ 34
Db 321 ELQHLWHTLWELLGAAMRQ 339

RESULT 8
E8262
ABC transporter ATP-binding protein XF1602 [imported] - Xylella fastidiosa (strain 9a5c)
C;Species: Xylella fastidiosa
C;Date: 18-Aug-2000 #sequence_revision 20-Aug-2000 #text_change 09-Jul-2004
C;Accession: E8262
R;Anonymous, The Xylella fastidiosa Consortium of the Organization for Nucleotide Sequen
Nature 406, 151-157, 2000
A;Title: The genome sequence of the plant pathogen Xylella fastidiosa.
A;Reference number: A82515; MUID:20365717; PMID:10910347
A;Note: for a complete list of authors see reference number A59328 below
A;Accession: E8262
A;Status: preliminary
A;Molecule type: DNA
A;Residues: 1-283 <SIM>
A;Cross-references: UNIPROT:Q9PD02; GB:AE003987; GB:AE003849; NID:G9106634; PDB:1AF8441
A;Experimental source: strain 9a5c
R;Simpon, A.U.G.; Reimach, F.C.; Arruda, P.; Abreu, F.A.; Acencio, M.; Alvarenga, R.; A
Birones, M.R.S.; Bueno, M.R.P.; Camargo, A.A.; Camargo, L.E.A.; Carraro, D.M.; Carier, H
as-Neto, E.; Docena, C.; El-Dorty, H.; Facincani, A.P.; Ferreira, A.J.S.
submitted to Genbank, June 2000
A;Authors: Ferreira, V.C.A.; Ferro, J.A.; Fraga, J.S.; Franco, S.C.; Franco, M.C.; From
J.D.; Junqueira, M.L.; Kemper, E.L.; Kitejima, J.P.; Krieger, J.E.; Kurame, E.B.; Laizy
Chado, M.A.; Madeira, A.M.B.N.; Madefra, H.M.F.; Martino, C.L.; Marques, M.V.; Martins, E
A;Authors: Martins, E.M.F.; Matsushima, A.Y.; Menck, C.F.M.; Miracca, E.C.; Miyaki, C.Y.;
F.G.; Nunes, L.R.; Oliveira, M.A.; de Oliveira, M.C.; de Oliveira, R.C.; Palmieri, D.A.
Rodrigues, V.; Rosa, A.J. de M.; de Rosa Jr., V.E.; de Sa, R.G.; Santelli, R.V.; Sawash

A;Authors: da Silva, A.C.R.; da Silva, F.R.; da Silva, A.M.; Silva Jr., W.A.; da Silveir
M.; Tshahko, M.H.; Vallada, H.; Van Sluys, M.A.; Verjovski-Almeida, S.; Vettore, A.L.; Z
A;Reference number: A59328
A;Contents: annotation
C;Genetics:
A;Gene: XF1602

Query Match 21.3%; Score 54; DB 2; Length 283;
Best Local Similarity 34.1%; Pred. No. 17;
Matches 15; Conservative 7; Mismatches 18; Indels 4; Gaps 1;

Qy 1 TQAQLRVGCVLTGTCOVONLSHRLMQLMGPRQDSAPVDPSSPSHY 40
Db 195 TDVAFIRDHGLVINTDVONLSHRLMQLMGPRQDSAPVDPSSPSHY 238

RESULT 9
T26050
hypothetical protein W01C.3 - Caenorhabditis elegans
C;Species: Caenorhabditis elegans
C;Date: 15-Oct-1999 #sequence_revision 15-Oct-1999 #text_change 09-Jul-2004
C;Accession: T26050
R;Sims, M.
submitted to the EMBL Data Library, June 1995
A;Reference number: Z20143
A;Accession: T26050
A;Status: preliminary; translated from GB/EMBL/DBJ
A;Molecule type: DNA
A;Residues: 1-792 <WIL>
A;Cross-references: UNIPROT:Q23115; EMBL:Z49969; PDB:1CAA90267.1; GSPDB:GN00020; CESP:W0
A;Experimental source: clone W01C9
C;Genetics:
A;Gene: CESP:W01C9.3
A;Map position: 2
A;Intons: 15/1; 40/1; 66/3; 101/3; 182/2; 270/3; 338/3; 366/1; 597/3; 613/2; 646/2; 684,

Query Match 21.3%; Score 54; DB 2; Length 792;
Best Local Similarity 39.5%; Pred. No. 50;
Matches 15; Conservative 3; Mismatches 18; Indels 2; Gaps 1;

Qy 9 GCVLGTCOVONLSHRLMQLMGPRQDSAPVDPSSPSHY 44
Db 736 GLALENRDADGFSHRLQQLMGTSNRYRDPVDPVNVNP 773

RESULT 10
A43708
gamma-interferon-inducible protein IP-30 precursor - human
C;Species: Homo sapiens (man)
C;Date: 03-Mar-1993 #sequence_revision 03-Mar-1993 #text_change 09-Jul-2004
C;Accession: A43708
R;Luster, A.D.; Weinshank, R.L.; Feilman, R.; Ravetch, J.V.
J. Biol. Chem. 263, 12036-12043, 1988
A;Title: Molecular and biochemical characterization of a novel gamma-interferon-inducible
A;Reference number: A43708; MUID:88298888; PMID:3136170
A;Accession: A43708
A;Status: preliminary
A;Molecule type: mRNA
A;Residues: 1-303 <LUS>
A;Cross-references: UNIPROT:P13284; EMBL:J03909; NID:G186264; PDB:1AAA36105.1; PID:G30704

Query Match 21.1%; Score 53.5; DB 2; Length 303;
Best Local Similarity 41.4%; Pred. No. 21;
Matches 12; Conservative 3; Mismatches 11; Indels 3; Gaps 1;

Qy 10 CVLGTGVONLSHRLMQLMGPRQDSAP 38
Db 209 CSPATARVCAIGHRWETL---GRSDPAP 234

RESULT 11
A37845
carboxylesterase (EC 3.1.1.1) precursor - Streptomyces scabies

C:Species: Streptomyces scabies
C:Date: 28-Jun-1991 #sequence_revision 28-Jun-1991 #text_change 09-Jul-2004
C:Accession: A37845; PGI103
R:Raymer, G.; Willard, J.M.A.; Schottel, J.L.
J. Bacteriol. 172, 7020-7026, 1990
A:Title: Cloning, sequencing, and regulation of expression of an extracellular esterase
A:Reference number: A37845; MUID:91072254; PMID:2254271
C:Accession: A37845
A:Molecule type: DNA
A:Residues: 1-345 <RAY>
A:Cross-references: UNIPROT:P22266; GB:M57297; NID:g153254; PIDN:AAA26743.1; PID:g153255
R:Schottel, J.L.; Hale, V.; Babcock, M.J.
Gene 115, 27-31, 1992
A:Title: Regulation and secretion of an extracellular esterase from Streptomyces scabies
A:Accession: PGI103; MUID:92307438; PMID:1612447
A:Reference number: PGI103
A:Molecule type: DNA
A:Residues: 1-54 <SCH>
A:Cross-references: GB:M57297
C:Comment: This protein is involved in suberin degradation.
C:Genetics:
A:Gene: est
C:Keywords: carboxylic ester hydrolase
F:1-35/Domain: signal sequence #status predicted <SIG>
F:40-345/Product: carboxylesterase #status predicted <MAT>

Query Match 21.1%; Score 53.5; DB 2; Length 345;
Best Local Similarity 34.2%; Pred. No. 24;
Matches 13; Conservative 3; Mismatches 13; Indels 9; Gaps 1;

4 QLRVGVLTGTCOVONLSHRLMQLMGPAGRODSAPVDP 41
16 RLTAALVAVLMSGLA-----LNGPAGSAGAPAPDP 44

RESULT 12
T35335
hypothetical protein SCSH.19c - Streptomyces coelicolor
C:Species: Streptomyces coelicolor
C:Date: 05-Nov-1999 #sequence_revision 05-Nov-1999 #text_change 09-Jul-2004
A:Accession: T35335
R:Oliver, K.; Harris, D.; James, K.D.; Parhill, J.; Barrell, B.G.; Rajandream, M.A.
submitted to the EMBL Data Library, May 1999
A:Reference number: Z21575
A:Accession: T35335
A:Status: preliminary; translated from GB/EMBL/DBJ
A:Molecule type: DNA
A:Residues: 1-136 <OLI>
A:Cross-references: UNIPROT:Q9X7S7; EMBL:AL049863; PIDN:CA842944.1; GSPDB:GN00070; SCOEI
A:Experimental source: strain A3(2)
C:Genetics:
A:Gene: SCOEDB:SCSH.19c

Query Match 20.9%; Score 53; DB 2; Length 136;
Best Local Similarity 38.1%; Pred. No. 10;
Matches 16; Conservative 3; Mismatches 19; Indels 4; Gaps 1;

7 RYGVLTGTCOVONLSHRLMQL-----MGPGRODSAPVDPSSP 44
19 RIGVPLGTAPRALALMLWRDRAWIAERLDPOEDPHLP 60

RESULT 13
AE2414
hypothetical protein all4869 [imported] - Nostoc sp. (strain PCC 7120)
C:Species: Nostoc sp. PCC 7120
A:Note: Nostoc sp. strain PCC 7120 is a synonym of Anabaena sp. strain PCC 7120
C:Date: 14-Dec-2001 #sequence_revision 14-Dec-2001 #text_change 09-Jul-2004
C:Accession: AE2414
R:Kaneko, T.; Nakamura, Y.; Wolk, C.P.; Kuritz, T.; Sasamoto, S.; Watanabe, A.; Iriguchi, N.; Shimpo, S.; Sugimoto, M.; Takazawa, M.; Yamada, M.; Yasuda, M.; Tabata, S.
DNA Res. 8, 205-213, 2001
A:Title: Complete Genomic Sequence of the Filamentous Nitrogen-fixing Cyanobacterium Ana

A:Reference number: AB1807; MUID:21595285; PMID:11759840
A:Accession: AE2414
A:Status: preliminary
A:Molecule type: DNA
A:Residues: 1-192 <KUR>
A:Cross-references: UNIPROT:Q8VNR4; GB:BA000019; PIDN:BAV6568.1; PID:g17134006; GSPDB:GN
A:Experimental source: strain PCC 7120
C:Genetics:
A:Gene: all4869
C:Superfamily: Synchocystis hypothetical protein slr1160

Query Match 20.9%; Score 53; DB 2; Length 192;
Best Local Similarity 40.0%; Pred. No. 15;
Matches 8; Conservative 6; Mismatches 6; Indels 0; Gaps 0;

27 LMGPAGRODSAPVDPSSPHS 46
23 LIAPVNAQNPPIDPNSPN 42

RESULT 14
H95954
probable Sir2-like transcription silencer protein [imported] - Sinorhizobium meliloti (str
C:Species: Sinorhizobium meliloti
C:Date: 24-Aug-2001 #sequence_revision 24-Aug-2001 #text_change 09-Jul-2004
C:Accession: H95954
R:Finan, T.M.; Weidner, S.; Wong, K.; Buhmester, J.; Chain, P.; Vorholter, F.J.; Hernan
Proc. Natl. Acad. Sci. U.S.A. 98, 9889-9894, 2001
A:Title: The complete sequence of the 1,663-bp pSymB megaplasmid from the N2-fixing endo
A:Reference number: A95842; MUID:21396508; PMID:11481431
A:Accession: H95954
A:Status: preliminary
A:Molecule type: DNA
A:Residues: 1-272 <KUR>
A:Cross-references: UNIPROT:P96452; GB:AL591985; PIDN:CA9304.1; PID:g15140790; GSPDB:GN
A:Experimental source: strain 1021, megaplasmid pSymB
R:Galibert, F.; Finan, T.M.; Long, S.R.; Puhler, A.; Abola, P.; Ampe, F.; Barloy-Hubler,
peta, D.; Chain, P.; Cowie, A.; Davis, R.W.; Dreano, S.; Federspiel, N.A.; Fisher, R.F.;
L.; Hyman, R.W.; Jones, T.
Science 293, 666-672, 2001
A:Author: Kahn, D.; Kahn, M.L.; Kallman, S.; Keating, D.H.; Kiss, E.; Komp, C.; Lelaure,
habault, P.; Vandendol, M.; Vorholter, F.J.; Weidner, S.; Wells, D.H.; Wong, K.; Yeh, K.C.
A:Title: The composite genome of the legume symbiont Sinorhizobium meliloti.
A:Reference number: A96039; MUID:21368234; PMID:11474104
A:Contents: annotation
C:Genetics:
A:Gene: Smb21328
A:Genome: plasmid
C:Superfamily: uncharacterized conserved protein with Sir2 domain

Query Match 20.9%; Score 53; DB 2; Length 272;
Best Local Similarity 26.1%; Pred. No. 22;
Matches 12; Conservative 10; Mismatches 22; Indels 2; Gaps 1;

2 QQLRVGVLTGTCOVONLSHRLMQLMGPAGRODSAPVDPSSPHSY 47
167 EATVPLGTGMSDNLIRLLIRIMWSIRSGHEKDRP--PEVVFMY 230

RESULT 15
E97187
dUDP-D-glucose 4,6-dehydratase [imported] - Clostridium acetobutylicum
C:Species: Clostridium acetobutylicum
C:Date: 14-Sep-2001 #sequence_revision 14-Sep-2001 #text_change 09-Jul-2004
C:Accession: E97187
R:Nolling, J.; Breton, G.; Omelchenko, M.V.; Markarova, K.S.; Zeng, Q.; Gibson, R.; Lee,
J.; Daly, M.J.; Bennett, G.N.; Kocin, E.V.; Smith, D.R.
J. Bacteriol. 183, 4823-4838, 2001
A:Title: Genome Sequence and Comparative Analysis of the Solvent-Producing Bacterium Clo
A:Reference number: A96900; MUID:21359325; PMID:21359325
A:Accession: E97187
A:Status: preliminary
A:Molecule type: DNA

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OM protein - protein search, using sw model

Run on: May 4, 2005, 18:06:10 ; Search time 41 Seconds
(without alignments)
85.573 Million cell updates/sec

Title: US-10-723-368-5
Perfect score: 254
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Scoring table: BIOSIM62
Gapop 10.0 , Gapect 0.5

Searched: 513545 seqs, 74649064 residues

Total number of hits satisfying chosen parameters: 513545

Minimum DB seq length: 0
Maximum DB seq length: 200000000

Post-processing: Minimum Match 0%

Listing first 45 summaries

Database : Issued Patents AA:*

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- 2: /cgn2_6/ptodata/1/1aa/5B_COMB.pep:*
- 3: /cgn2_6/ptodata/1/1aa/6A_COMB.pep:*
- 4: /cgn2_6/ptodata/1/1aa/6B_COMB.pep:*
- 5: /cgn2_6/ptodata/1/1aa/PTUS_COMB.pep:*
- 6: /cgn2_6/ptodata/1/1aa/backfile1.pep:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	72.5	28.5	38	4	US-09-280-501-6
2	72.5	28.5	40	4	US-09-280-501-11
3	72.5	28.5	52	3	US-09-070-504-14
4	72.5	28.5	52	4	US-09-813-345C-14
5	72.5	28.5	185	1	US-08-233-389C-1
6	72.5	28.5	185	2	US-08-801-863-1
7	72.5	28.5	185	2	US-08-486-596A-1
8	72.5	28.5	185	2	US-09-004-713-1
9	70.5	27.8	188	1	US-08-233-389C-3
10	70.5	27.8	188	2	US-08-801-863-3
11	70.5	27.8	188	2	US-08-486-596A-3
12	70.5	27.8	188	2	US-09-004-713-3
13	69.5	27.4	50	3	US-09-070-504-15
14	69.5	27.4	50	4	US-09-813-345C-15
15	65.5	25.8	40	4	US-09-280-501-8
16	65.5	25.8	50	4	US-09-280-501-7
17	58	22.8	559	2	US-08-406-855A-20
18	58	22.8	559	3	US-09-206-899-20
19	58	22.8	560	4	US-09-688-415-8
20	56	22.0	172	4	US-09-949-016-9374
21	56	22.0	770	4	US-09-252-991A-28510
22	55.5	21.9	906	4	US-09-252-991A-28132
23	54.5	21.5	184	4	US-09-252-991A-32339
24	54	21.3	133	4	US-09-252-991A-30594
25	53.5	21.1	218	4	US-09-270-767-43455
26	53.5	21.1	303	4	US-09-949-016-6717
27	53	20.9	181	4	US-09-530-685A-21

28	53	20.9	239	4	US-09-530-685A-32	Sequence 32, Appl
29	53	20.9	255	4	US-09-902-540-14178	Sequence 14178, A
30	53	20.9	309	4	US-09-530-685A-31	Sequence 31, Appl
31	53	20.9	450	3	US-09-120-365-68	Sequence 68, Appl
32	53	20.9	450	3	US-09-515-039-68	Sequence 68, Appl
33	53	20.9	605	3	US-08-753-007A-2	Sequence 2, Appl
34	53	20.9	605	3	US-09-398-496-2	Sequence 2, Appl
35	53	20.9	754	2	US-08-525-864A-2	Sequence 2, Appl
36	52	20.5	263	4	US-09-270-767-46206	Sequence 46206, A
37	52	20.5	675	4	US-09-252-991A-32681	Sequence 32681, A
38	52	20.5	3898	3	US-08-750-717-2	Sequence 2, Appl
39	51.5	20.3	149	4	US-09-543-681A-8072	Sequence 8072, Ap
40	51.5	20.3	155	4	US-09-252-991A-20281	Sequence 20281, A
41	51.5	20.3	220	4	US-09-902-540-11202	Sequence 11202, A
42	51.5	20.3	879	4	US-09-252-991A-31990	Sequence 31990, A
43	51	20.1	1213	2	US-08-937-102-2	Sequence 2, Appl
44	51	20.1	1399	4	US-09-388-221B-4	Sequence 4, Appl
45	51	20.1	1424	4	US-09-388-221B-12	Sequence 12, Appl

ALIGNMENTS

```
RESULT 1
US-09-280-501-6
Sequence 6, Application US/09280501
Patent No. 6440421
GENERAL INFORMATION:
APPLICANT: Cooper, Garth James Smith
APPLICANT: Reid, Ian Reginald
TITLE OF INVENTION: TREATMENT OF BONE DISORDERS WITH
ADRENOMEDULLIN OR ADRENOMEDULLIN AGONISTS
FILE REFERENCE: 08987-005001
CURRENT APPLICATION NUMBER: US/09/280,501
PRIOR FILING DATE: 1999-03-30
PRIOR APPLICATION NUMBER: 08/634,562
NUMBER OF SEQ ID NOS: 17
SOFTWARE: FastSeq for Windows Version 4.0
SEQ ID NO 6
LENGTH: 38
TYPE: PRT
ORGANISM: Homo sapiens
US-09-280-501-6

Query Match 28.5%; Score 72.5; DB 4; Length 38;
Best Local Similarity 38.5%; Pred. No. 0.0098;
Matches 15; Conservative 6; Mismatches 17; Indels 1;

Cy 9 GCVLTGTCVQVNLSHRLMQLMGPRGRDSAPVDPSSPHSY 47
Db 1 GCRFGTCTVOKLHAQIYQFT-DKDKDVAPRSKISPGY 38

RESULT 2
US-09-280-501-11
Sequence 11, Application US/09280501
Patent No. 6440421
GENERAL INFORMATION:
APPLICANT: Cooper, Garth James Smith
APPLICANT: Reid, Ian Reginald
TITLE OF INVENTION: TREATMENT OF BONE DISORDERS WITH
ADRENOMEDULLIN OR ADRENOMEDULLIN AGONISTS
FILE REFERENCE: 08987-005001
CURRENT APPLICATION NUMBER: US/09/280,501
PRIOR FILING DATE: 1999-03-30
PRIOR APPLICATION NUMBER: 08/634,562
NUMBER OF SEQ ID NOS: 17
SOFTWARE: FastSeq for Windows Version 4.0
SEQ ID NO 11
```

LENGTH: 40
TYPE: PRT
ORGANISM: Homo sapiens
US-09-280-501-11

Query Match 28.5%; Score 72.5; DB 4; Length 40;
Best Local Similarity 38.5%; Pred. No. 0.01;
Matches 15; Conservative 6; Mismatches 17; Indels 1; Gaps 1;

QY 9 GCVLGTCQVONLSHRLMQLMGPAQRDSAPVDPSSPHSY 47
DB 3 GCRFGTCVQKLAHQIYQFT-DKDKDNVAPRSKISPOGY 40

RESULT 3

US-09-070-504-14
Sequence 14, Application US/09070504
Patent No. 6268474

GENERAL INFORMATION:
APPLICANT: Smith, Derek D.
APPLICANT: Saha, Shankar
APPLICANT: Abel, Peter W.
TITLE OF INVENTION: PEPTIDE ANTAGONISTS OF GCRP-RECEPTOR
TITLE OF INVENTION: SUPERFAMILY AND METHODS OF USE
NUMBER OF SEQUENCES: 23
CORRESPONDENCE ADDRESS:
ADDRESSEE: Muehling, Raasch & Gebhardt, P.A.
STREET: 119 No. 6268474th Fourth Street
CITY: Minneapolis
STATE: MN
COUNTRY: USA
ZIP: 55401

COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/070,504
FILING DATE: 30-APR-1998
CLASSIFICATION:
ATTORNEY/AGENT INFORMATION:
NAME: McCormack, Myra H
REGISTRATION NUMBER: 36,602
REFERENCE/DOCKET NUMBER: 180.00020101
TELECOMMUNICATION INFORMATION:
TELEPHONE: 612/305-1220
TELEFAX: 612/305-1228
INFORMATION FOR SEQ ID NO: 14:
SEQUENCE CHARACTERISTICS:
LENGTH: 52 amino acids
TYPE: amino acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: peptide

US-09-070-504-14

Query Match 28.5%; Score 72.5; DB 3; Length 52;
Best Local Similarity 38.5%; Pred. No. 0.014;
Matches 15; Conservative 6; Mismatches 17; Indels 1; Gaps 1;

QY 9 GCVLGTCQVONLSHRLMQLMGPAQRDSAPVDPSSPHSY 47
DB 15 GCRFGTCVQKLAHQIYQFT-DKDKDNVAPRSKISPOGY 52

RESULT 4
US-09-813-345C-14
Sequence 14, Application US/09813345C
Patent No. 6756205

GENERAL INFORMATION:
APPLICANT: CREIGHTON UNIVERSITY
APPLICANT: SMITH, Derek D.

APPLICANT: SAHA, Shankar
APPLICANT: ABEL, Peter W.
TITLE OF INVENTION: PEPTIDE ANTAGONISTS OF GCRP-RECEPTOR SUPERFAMILY AND METHODS OF
TITLE OF INVENTION: USE
FILE REFERENCE: 180.00020102
CURRENT APPLICATION NUMBER: US/09/813,345C
CURRENT FILING DATE: 2001-03-20
PRIOR APPLICATION NUMBER: 09/070,504
PRIOR FILING DATE: 1998-04-30
NUMBER OF SEQ ID NOS: 23
SOFTWARE: Patentin version 3.2
SEQ ID NO 14
LENGTH: 52
TYPE: PRT
ORGANISM: Artificial Sequence
OTHER INFORMATION: Artificially Synthesized Peptide
US-09-813-345C-14

Query Match 28.5%; Score 72.5; DB 4; Length 52;
Best Local Similarity 38.5%; Pred. No. 0.014;
Matches 15; Conservative 6; Mismatches 17; Indels 1; Gaps 1;

QY 9 GCVLGTCQVONLSHRLMQLMGPAQRDSAPVDPSSPHSY 47
DB 15 GCRFGTCVQKLAHQIYQFT-DKDKDNVAPRSKISPOGY 52

RESULT 5

US-08-233-389C-1
Sequence 1, Application US/08233389C
Patent No. 5639855

GENERAL INFORMATION:
APPLICANT: KITAMURA, Kazuo
APPLICANT: KANGAWA, Kenji
APPLICANT: MATSUDO, Hisayuki
APPLICANT: ETO, Tanenao
TITLE OF INVENTION: ADRENOMEDULLIN
NUMBER OF SEQUENCES: 10
CORRESPONDENCE ADDRESS:
ADDRESSEE: C/O FISH & NEAVE
STREET: 1251 Avenue of the Americas
CITY: New York
STATE: NY
COUNTRY: USA
ZIP: 10020

COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/233,389C
FILING DATE: 26-APR-1994
CLASSIFICATION: 530
ATTORNEY/AGENT INFORMATION:
NAME: HALEY Jr., James F.
REGISTRATION NUMBER: 27,794
REFERENCE/DOCKET NUMBER: SHGN-5
TELECOMMUNICATION INFORMATION:
TELEPHONE: (212) 596-9000
TELEFAX: (212) 596-9090
INFORMATION FOR SEQ ID NO: 1:
SEQUENCE CHARACTERISTICS:
LENGTH: 185 amino acids
TYPE: amino acid
TOPOLOGY: linear
MOLECULE TYPE: protein

US-08-233-389C-1

Query Match 28.5%; Score 72.5; DB 1; Length 185;
Best Local Similarity 38.5%; Pred. No. 0.056;
Matches 15; Conservative 6; Mismatches 17; Indels 1; Gaps 1;

Qy 9 GCVLGTQOVONLSHRLWQMGAPAGODSAPVDPSSPSHY 47
Db 109 GCRFGTCTVQKLAHQIYQFT-DKDKXNVAPRSKISPGCY 146

RESULT 6

US-08-801-863-1
Sequence 1, Application US/08801863
Patent No. 5830703
GENERAL INFORMATION:
APPLICANT: KITAMURA, Kazuo
APPLICANT: KANGAWA, Kenji
APPLICANT: MATSUO, Hisayuki
APPLICANT: ETO, Tanenao
TITLE OF INVENTION: ADRENOMEDULLIN
NUMBER OF SEQUENCES: 10
CORRESPONDENCE ADDRESS:
ADDRESSEE: c/o FISH & NEAVE
STREET: 1251 Avenue of the Americas
CITY: New York
STATE: NY
COUNTRY: USA
ZIP: 10020
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patent in Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/801,863
FILING DATE: CONCURRENTLY HERewith
CLASSIFICATION: 530
ATTORNEY/AGENT INFORMATION:
NAME: HALEY Jr., James F.
REGISTRATION NUMBER: 27,794
REFERENCE/DOCKET NUMBER: SHGN-5 DIV3
TELECOMMUNICATION INFORMATION:
TELEPHONE: (212) 596-9000
TELEFAX: (212) 596-9090
INFORMATION FOR SEQ ID NO: 1:
SEQUENCE CHARACTERISTICS:
LENGTH: 185 amino acids
TYPE: amino acid
TOPOLOGY: linear
MOLECULE TYPE: protein
US-08-801-863-1

Query Match 28.5%; Score 72.5; DB 2; Length 185;
Best Local Similarity 38.5%; Pred. No. 0.056; Matches 15; Conservative 6; Mismatches 17; Indels 1; Gaps 1;

Qy 9 GCVLGTQOVONLSHRLWQMGAPAGODSAPVDPSSPSHY 47
Db 109 GCRFGTCTVQKLAHQIYQFT-DKDKXNVAPRSKISPGCY 146

RESULT 7

US-08-486-596A-1
Sequence 1, Application US/08486596A
Patent No. 5837823
GENERAL INFORMATION:
APPLICANT: KITAMURA, Kazuo
APPLICANT: KANGAWA, Kenji
APPLICANT: MATSUO, Hisayuki
APPLICANT: ETO, Tanenao
TITLE OF INVENTION: ADRENOMEDULLIN
NUMBER OF SEQUENCES: 10
CORRESPONDENCE ADDRESS:
ADDRESSEE: c/o FISH & NEAVE
STREET: 1251 Avenue of the Americas
CITY: New York
STATE: NY

COUNTRY: USA
ZIP: 10020
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patent in Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/486,596A
FILING DATE: JUNE 7, 1995
CLASSIFICATION: 530
ATTORNEY/AGENT INFORMATION:
NAME: HALEY Jr., James F.
REGISTRATION NUMBER: 27,794
REFERENCE/DOCKET NUMBER: SHGN-5 DIV1
TELECOMMUNICATION INFORMATION:
TELEPHONE: (212) 596-9000
TELEFAX: (212) 596-9090
INFORMATION FOR SEQ ID NO: 1:
SEQUENCE CHARACTERISTICS:
LENGTH: 185 amino acids
TYPE: amino acid
TOPOLOGY: linear
MOLECULE TYPE: protein
US-08-486-596A-1

Query Match 28.5%; Score 72.5; DB 2; Length 185;
Best Local Similarity 38.5%; Pred. No. 0.056; Matches 15; Conservative 6; Mismatches 17; Indels 1; Gaps 1;

Qy 9 GCVLGTQOVONLSHRLWQMGAPAGODSAPVDPSSPSHY 47
Db 109 GCRFGTCTVQKLAHQIYQFT-DKDKXNVAPRSKISPGCY 146

RESULT 8

US-09-004-713-1
Sequence 1, Application US/09004713
Patent No. 5910416
GENERAL INFORMATION:
APPLICANT: KITAMURA, Kazuo
APPLICANT: KANGAWA, Kenji
APPLICANT: MATSUO, Hisayuki
APPLICANT: ETO, Tanenao
TITLE OF INVENTION: ADRENOMEDULLIN
NUMBER OF SEQUENCES: 10
CORRESPONDENCE ADDRESS:
ADDRESSEE: c/o FISH & NEAVE
STREET: 1251 Avenue of the Americas
CITY: New York
STATE: NY
COUNTRY: USA
ZIP: 10020
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patent in Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/004,713
FILING DATE: JANUARY 7, 1998
CLASSIFICATION: 435
ATTORNEY/AGENT INFORMATION:
NAME: HALEY Jr., James F.
REGISTRATION NUMBER: 27,794
REFERENCE/DOCKET NUMBER: SHGN-5 DIV2 CON
TELECOMMUNICATION INFORMATION:
TELEPHONE: (212) 596-9000
TELEFAX: (212) 596-9090
INFORMATION FOR SEQ ID NO: 1:
SEQUENCE CHARACTERISTICS:
LENGTH: 185 amino acids
TYPE: amino acid

TOPOLOGY: linear
US-09-004-713-1

Query Match
Best Local Similarity 28.5%; Score 72.5; DB 2; Length 185;
Matches 15; Conservative 6; Mismatches 17; Indels 1; Gaps 1;
Qy 9 GCVLGTQVONLSHRLMQLMGPAGRODSAPVDSSPSHY 47
Db 109 GCRFGTCTVOKLAHQIYQFT-DKDKDGVAPRSKISFGY 146

RESULT 9
US-08-233-389C-3
Sequence 3, Application US/08233389C
Patent No. 5639655
GENERAL INFORMATION:
APPLICANT: KITAMURA, Kazuo
APPLICANT: KANGAWA, Kenji
APPLICANT: MATSUO, Hisayuki
TITLE OF INVENTION: ADRENOMEDULLIN
NUMBER OF SEQUENCES: 10
CORRESPONDENCE ADDRESS:
ADDRESSEE: C/O FISH & NEAVE
STREET: 1251 Avenue of the Americas
CITY: New York
STATE: NY
COUNTRY: USA
ZIP: 10020
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/233.389C
FILING DATE: 26-APR-1994
CLASSIFICATION: 530
ATTORNEY/AGENT INFORMATION:
NAME: HALEY Jr., James F.
REGISTRATION NUMBER: 27,794
REFERENCE/DOCKET NUMBER: SHGN-5
TELECOMMUNICATION INFORMATION:
TELEPHONE: (212) 596-9000
TELEFAX: (212) 596-9000
INFORMATION FOR SEQ ID NO: 3:
SEQUENCE CHARACTERISTICS:
LENGTH: 188 amino acids
TYPE: amino acid
TOPOLOGY: linear
MOLECULE TYPE: protein
US-08-233-389C-3

Query Match
Best Local Similarity 27.8%; Score 70.5; DB 1; Length 186;
Matches 15; Conservative 5; Mismatches 18; Indels 1; Gaps 1;
Qy 9 GCVLGTQVONLSHRLMQLMGPAGRODSAPVDSSPSHY 47
Db 109 GCRFGTCTVOKLAHQIYQFT-DKDKDGVAPRSKISFGY 146

RESULT 10
US-08-801-863-3
Sequence 3, Application US/08801863
Patent No. 5830703
GENERAL INFORMATION:
APPLICANT: KITAMURA, Kazuo
APPLICANT: KANGAWA, Kenji
APPLICANT: MATSUO, Hisayuki
APPLICANT: ETO, Tanenao

TITLE OF INVENTION: ADRENOMEDULLIN
NUMBER OF SEQUENCES: 10
CORRESPONDENCE ADDRESS:
ADDRESSEE: C/O FISH & NEAVE
STREET: 1251 Avenue of the Americas
CITY: New York
STATE: NY
COUNTRY: USA
ZIP: 10020
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/801.863
FILING DATE: JUNE 7, 1995
CLASSIFICATION: 530
ATTORNEY/AGENT INFORMATION:
NAME: HALEY Jr., James F.
REGISTRATION NUMBER: 27,794
REFERENCE/DOCKET NUMBER: SHGN-5 DIV3
TELECOMMUNICATION INFORMATION:
TELEPHONE: (212) 596-9000
TELEFAX: (212) 596-9000
INFORMATION FOR SEQ ID NO: 3:
SEQUENCE CHARACTERISTICS:
LENGTH: 188 amino acids
TYPE: amino acid
TOPOLOGY: linear
MOLECULE TYPE: protein
US-08-801-863-3

Query Match
Best Local Similarity 27.8%; Score 70.5; DB 2; Length 186;
Matches 15; Conservative 5; Mismatches 18; Indels 1; Gaps 1;
Qy 9 GCVLGTQVONLSHRLMQLMGPAGRODSAPVDSSPSHY 47
Db 109 GCRFGTCTVOKLAHQIYQFT-DKDKDGVAPRSKISFGY 146

RESULT 11
US-08-486-596A-3
Sequence 3, Application US/08486596A
Patent No. 5837823
GENERAL INFORMATION:
APPLICANT: KITAMURA, Kazuo
APPLICANT: KANGAWA, Kenji
APPLICANT: MATSUO, Hisayuki
TITLE OF INVENTION: ADRENOMEDULLIN
NUMBER OF SEQUENCES: 10
CORRESPONDENCE ADDRESS:
ADDRESSEE: C/O FISH & NEAVE
STREET: 1251 Avenue of the Americas
CITY: New York
STATE: NY
COUNTRY: USA
ZIP: 10020
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/486.596A
FILING DATE: JUNE 7, 1995
CLASSIFICATION: 530
ATTORNEY/AGENT INFORMATION:
NAME: HALEY Jr., James F.
REGISTRATION NUMBER: 27,794
REFERENCE/DOCKET NUMBER: SHGN-5 DIV1

TELECOMMUNICATION INFORMATION:
TELEPHONE: (212) 596-9000
TELEFAX: (212) 596-9090
INFORMATION FOR SEQ ID NO: 3:
SEQUENCE CHARACTERISTICS:
LENGTH: 188 amino acids
TYPE: amino acid
TOPOLOGY: linear
MOLECULE TYPE: protein
US-08-486-596A-3

Query Match 27.8%; Score 70.5; DB 2; Length 188;
Best Local Similarity 38.5%; Pred. No. 0.11;
Matches 15; Conservative 5; Mismatches 18; Indels 1; Gaps 1;

Qy 9 GCVLGTQVONLSHRLMQLMGPAQRDSAPVDPSSPHSY 47
Db 109 GCRFGTCTVQKLAHQIYQFT-DKDKGAVAPRSKISPGGY 146

RESULT 12
US-09-004-713-3
Sequence 3, Application US/09004713
Patent No. 5910416

GENERAL INFORMATION:
APPLICANT: KITAMURA, Kazuo
APPLICANT: KANAGAWA, Kenji
APPLICANT: MATSUO, Hisayuki
APPLICANT: ETO, Tanemao
TITLE OF INVENTION: ADRENOMEDULLIN
NUMBER OF SEQUENCES: 10
CORRESPONDENCE ADDRESS:
ADDRESSEE: C/O FISH & NEAVE
STREET: 1251 Avenue of the Americas
CITY: New York
STATE: NY
COUNTRY: USA
ZIP: 10020

COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/004,713
FILING DATE: JANUARY 7, 1998
CLASSIFICATION: 435
ATTORNEY/AGENT INFORMATION:
NAME: HALEY Jr., James F.
REGISTRATION NUMBER: 27,794
REFERENCE/DOCKET NUMBER: SHGN-5 DIV2 CON
TELECOMMUNICATION INFORMATION:
TELEPHONE: (212) 596-9000
TELEFAX: (212) 596-9090
INFORMATION FOR SEQ ID NO: 3:
SEQUENCE CHARACTERISTICS:
LENGTH: 188 amino acids
TYPE: amino acid
TOPOLOGY: linear
MOLECULE TYPE: protein
US-09-004-713-3

Query Match 27.8%; Score 70.5; DB 2; Length 188;
Best Local Similarity 38.5%; Pred. No. 0.11;
Matches 15; Conservative 5; Mismatches 18; Indels 1; Gaps 1;

Qy 9 GCVLGTQVONLSHRLMQLMGPAQRDSAPVDPSSPHSY 47
Db 109 GCRFGTCTVQKLAHQIYQFT-DKDKGAVAPRSKISPGGY 146

RESULT 13
US-09-070-504-15

Sequence 15, Application US/09070504
Patent No. 6268474

GENERAL INFORMATION:
APPLICANT: Smith, Derek D.
APPLICANT: Saha, Shankar
APPLICANT: Abel, Peter W.
TITLE OF INVENTION: PEPTIDE ANTAGONISTS OF CGRP-RECEPTOR
TITLE OF INVENTION: SUPERFAMILY AND METHODS OF USE
NUMBER OF SEQUENCES: 23
CORRESPONDENCE ADDRESS:
ADDRESSEE: Muehling, Raasch & Gebhardt, P.A.
STREET: 119 No. 6268474th Fourth Street
CITY: Minneapolis
STATE: MN
COUNTRY: USA
ZIP: 55401

COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/070,504
FILING DATE: 30-Apr-1998
CLASSIFICATION:
ATTORNEY/AGENT INFORMATION:
NAME: McCormack, Myra H
REGISTRATION NUMBER: 36,602
REFERENCE/DOCKET NUMBER: 180.00020101
TELECOMMUNICATION INFORMATION:
TELEPHONE: 612/305-1220
TELEFAX: 612/305-1228
INFORMATION FOR SEQ ID NO: 15:
SEQUENCE CHARACTERISTICS:
LENGTH: 50 amino acids
TYPE: amino acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: peptide
US-09-070-504-15

Query Match 27.4%; Score 69.5; DB 3; Length 50;
Best Local Similarity 35.9%; Pred. No. 0.033;
Matches 14; Conservative 7; Mismatches 17; Indels 1; Gaps 1;

Qy 9 GCVLGTQVONLSHRLMQLMGPAQRDSAPVDPSSPHSY 47
Db 13 GCRFGTCTVQKLAHQIYQFT-DKDKGMAPRNKISPGGY 50

RESULT 14
US-09-813-345C-15
Sequence 15, Application US/09813345C
Patent No. 6756205

GENERAL INFORMATION:
APPLICANT: CREIGHTON UNIVERSITY
APPLICANT: SMITH, Derek D.
APPLICANT: SAHA, Shankar
APPLICANT: ABEL, Peter W.
TITLE OF INVENTION: PEPTIDE ANTAGONISTS OF CGRP-RECEPTOR SUPERFAMILY AND METHODS OF
TITLE OF INVENTION: USE
FILE REFERENCE: 180.00020102
CURRENT APPLICATION NUMBER: US/09/813,345C
CURRENT FILING DATE: 2001-03-20
PRIOR APPLICATION NUMBER: 09/070,504
PRIOR FILING DATE: 1998-04-30
NUMBER OF SEQ ID NOS: 23
SOFTWARE: PatentIn version 3.2
SEQ ID NO 15
LENGTH: 50
TYPE: PRT
ORGANISM: Artificial Sequence
FEATURE:

US-09-070-504-15

OTHER INFORMATION: Artificially Synthesized Peptide
US-09-813-345C-15

Query Match 27.4%; Score 69.5; DB 4; Length 50;
Best Local Similarity 35.9%; Pred. No. 0.033;
Matches 14; Conservative 7; Mismatches 17; Indels 1; Gaps 1;

OY 9 GCVLGTCOVONLSHRLMOLMGPAQRDSAPVDPSPHSY 47
DB 13 GCRFGCTVOKLAHQIYQFT-DKDKDMAPRNKISPGY 50

RESULT 15

US-09-280-501-8
; Sequence 8; Application US/09280501
; Patent No. 6440421
; GENERAL INFORMATION:
; APPLICANT: Cooper, Garth James Smith
; APPLICANT: Reid, Ian Reginald
; APPLICANT: Cornish, Jillian
; TITLE OF INVENTION: TREATMENT OF BONE DISORDERS WITH
; FILE REFERENCE: 08987-005001
; CURRENT FILING DATE: 1999-03-30
; PRIOR APPLICATION NUMBER: 08/534,562
; PRIOR FILING DATE: 1996-04-18
; NUMBER OF SEQ ID NOS: 17
; SOFTWARE: FASTSEQ for Windows Version 4.0
; SEQ ID NO 8
; LENGTH: 40
; TYPE: PRT
; ORGANISM: Homo sapiens
US-09-280-501-8

Query Match 25.8%; Score 65.5; DB 4; Length 40;
Best Local Similarity 38.9%; Pred. No. 0.089;
Matches 14; Conservative 6; Mismatches 15; Indels 1; Gaps 1;

OY 9 GCVLGTCOVONLSHRLMOLMGPAQRDSAPVDPSP 44
DB 5 GCRFGCTVOKLAHQIYQFT-DKDKDMAPRNKIS 39

Search completed: May 4, 2005, 18:15:51
Job time : 50 secs